

Prepared for The Sac & Fox Nation Of Missouri in Kansas and Nebraska

High-Resolution Light NonAqueous Phase Liquid (LNAPL) Conceptual Site Model Assessment, Sac & Fox Truck Stop, 1346 US Highway 75, Powhattan, Kansas March 2018

Project Number 3672-2017-10



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Conversion Factors

Inch/Ounce/Pound/PSI to International System of Units

Multiply	Ву	To obtain
	Length	
Inch (in.)	2.54	Centimeter (cm)
Inch (in.)	25.4	Millimeter (mm)
Foot (ft.)	0.3048	Meter (m)
	Volume	
Ounce (oz.)	29.6	Milliliters (ml)
Gallon (gal)	3.8	Liters (L)
	Pressure	
Pounds per Square Inch (psi)	6.89	Kilopascals (kPa)
	Hydraulic Conductivity	
Feet per day (ft/day)	0.0003527	Centimeters per second (cm/sec)

Temperature in degrees Celsius (°C) is converted to degrees Fahrenheit (°F) as

$$(^{\circ}F) = (1.8 \times (^{\circ}C) + 32)$$

Datum

Horizontal and vertical coordinates are referenced from the World Geodetic System 1984 [EPSG:4326].

Supplemental Information

Electrical conductivity (EC) is provided in millisiemens per meter (mS/meter).

Concentrations of chemical constituents in water are provided in either milligrams per liter (mg/L) or micrograms per liter (µg/L).

Concentrations of chemical constituents in soil are provided in either milligrams per kilogram (mg/kg) or micrograms per kilogram (μ g/kg).

Concentrations of chemical constituents in vapor are provided in either milligrams per cubic meter (mg/m^3) or micrograms per cubic meter ($\mu g/m^3$).

High-Resolution Light NonAqueous Phase Liquid (LNAPL) Conceptual Site Model Assessment, Sac & Fox Truck Stop, 1346 US Highway 75, Powhattan, Kansas March 2018

Summary

COLUMBIA Technologies, LLC, in collaboration with The Sac & Fox Nation Of Missouri in Kansas and Nebraska (Sac & Fox), conducted a high-resolution assessment of the Sac & Fox Truck Stop, 1346 US Highway 75, in Powhattan, Kansas (the Site) during the period of 19 to 23 March 2018.

The primary objectives of this assessment were to isolate and characterize the extent and distribution of LNAPL and to identify and describe any residual zones of petroleum acting as residual sources and any likely pathway(s) for preferential LNAPL migration to support the business goals of monetizing the long-term expense of LNAPL management. To accomplish this objective, a High-Resolution Site Characterization (HRSC) was conducted, and an updated LNAPL Conceptual Site Model (LCSM) was developed.

The updated LCSM provides a better understanding for:

- (1) evaluating the risks associated with the residual LNAPL remaining at the site;(2) setting achievable remedial action goals, and
- (3) selecting or modifying effective remedial action technologies.

This HRSC was conducted in accordance with the guidelines of the Interstate Training and Regulatory Council (ITRC), Evaluating LNAPL Remedial Technologies for Achieving Project Goals (Dec 2009).

The updated LCSM presented herein is based on high-resolution direct sensing measurements made by COLUMBIA

Technologies and pertinent historic site data provided by Sac & Fox and their primary consultant Terranext LLC. Historical indications of residual LNAPL are presented in Figure 1.

The direct sensing data employed for this assessment are comprised of Laser Induced Fluorescence/Ultraviolet Optical Screening Tool (LIF/UVOST®) and combined Membrane Interface Probe (MIP) and Hydraulic Profiling Tool (HPT) measurements. This combined tool is referred to as a MiHpt. Direct sensing logs are presented in **Appendices D**, **E**, and **F**.

Direct sensing survey stations are shown in **Figure 2**. Direct sensing survey locations consist of the following:

- Eight (8) LIF/UVOST® borings
- Fifteen (15) MiHpt borings

On March 23, 2018, following the direct sensing survey high-resolution soil sampling was systematically conducted at the Site at stations where LNAPL or high concentrations of petroleum hydrocarbons (PHCs) were identified with the direct sensing tools.

Twenty-six (26) discrete soil samples were collected from equally spaced vertical intervals through the apparent LNAPL or PHC horizon at five stations SF07, 11, 14, 16, and 18 using TerraCore® non-methanol soil samplers in accordance with EPA Method 5035. The soil samples were shipped to the ALS Environmental Laboratory to measure TPH concentrations in both the gasoline range (GRO) and diesel range (DRO) using Texas Method TX1005. These high-resolution soil sampling stations are also identified in **Figure 2**.

This high-resolution assessment of the **Sac & Fox Truck Stop** delineated residual LNAPL impacts remain at much of the site based on multiple lines of evidence that include direct sensing survey results, discrete soil sampling and analysis, and elevated BTEX concentrations in groundwater.

Gasoline range LNAPL was identified at the site covering a footprint from the northwest portion of the property eastward with residual impacts at the north, south, and eastern boundaries as shown in **Figure 3**.

Both LIF-UVOST® and soil analytical results in the C12-C28 range at station SF14 indicate the probable presence of diesel fuel at fourteen (14) to fifteen (15) feet bgs.

Residual LNAPL was measured in a vertical interval of five (5) to fifteen (15) feet below grade (bgs) on the upper operating portion of the property as shown in **Figures 4 through 8.**

Soil concentrations for TPH indicative of residual LNAPL are present within thirty (30) feet of the occupied buildings and more shallow than fifteen (15) feet below grade. These conditions exceed the preliminary screening for potential petroleum vapor

intrusion into the building. Previous soil vapor measurements taken in January 2017 at a depth of six (6) feet below grade were reported by Terranext. The results included two measurements of 489,000 ug/m³ at station SV-1 and 235,000 ug/m³ at station SV-2. As previously reported by Terranext LLC, these levels exceed those recommended by the U.S. EPA.

The primary source of the residual LNAPL appears to be from the vicinity of the former UST enclosure to the west of monitoring well MW-6.

Multiple lines of evidence developed through this HRSC are indicative of a residual, predominantly immobile, LNAPL source zone. This implies that the overall LNAPL "footprint" is stable on a macro-scale although localized LNAPL movement into and out of pore spaces (or monitoring wells) may persist.

Most monitoring wells are installed with screen intervals in soils exhibiting low hydraulic permeability. The tops of the monitoring well screens are at times below the reported air-water interface and the bulk of the residual petroleum hydrocarbon mass. Both of these factors will likely result in groundwater concentrations for petroleum hydrocarbons being biased low. Additionally, the effectiveness of the monitoring wells for any planned groundwater treatment are diminished.

Introduction

Background and Current Conceptual Site Model (Background information provided by Terranext LLC are in italics)

The Sac & Fox Truck Stop is located at **1346 US Highway 75**, in Powhattan, Kansas.

The Site is an operating retail gasoline station with concrete slab under the canopies, concrete over the tank basin, and remainder of the site is asphalt pavement.

On July 20, 2015, Sac & Fox reported a release of gasoline from an existing 15,000-gallon underground storage tank (UST). Due to heavy rains, approximately 63,000 gallons of impacted groundwater was pumped from the UST basin and LUST on July 10th -11th and 25th -26th, 2016.

Conclusions from the historical monitoring well installation activities, soil sampling, groundwater sampling, initial CSM development and initial soil vapor screening are as follows:

- Based on the soil sampling, subsurface soils have been impacted above KDHE's Tier 2 non-residential RSK values at the MW4 and MW6 locations.
- Based on the groundwater sampling, groundwater has been impacted above KDHE's Tier 2 non-residential RSK values at the MW-1, MW-3, MW-4 and MW-6 locations.
- Based on the groundwater gauging, groundwater flow near the Sac & Fox Truck Stop facility is generally towards the east, towards the surface pond located approximately 600 feet east-southeast of the facility.

Based on the monitoring well installation activities, there appears to be an orange silty sand and sand unit encountered approximately 12-20 feet bgs. This unit appears to become more prevalent to the east.

- Based on the initial soil vapor screening, there exists the potential for soil vapor intrusion into the existing facility building.
- Based on the initial CSM, there exist pathways for impacted subsurface soils and groundwater to reach receptors. Potentially impacted receptors included: an ecological receptor (the pond and associated animals) down gradient (east) of the facility; commercial works on-site, and construction workers should construction activities occur at the facility.

A soil vapor survey was performed by Terranext on January 24, 2017. Soil vapor data at near-source depth (6 feet below ground surface (bgs)) and sub-slab depth (3-feet bgs) at two locations on the south side of the existing building to determine potential vapor intrusion impacts. Soil vapor concentrations were determined to be well above screening levels calculated using U.S. EPA's MAY 2016 Vapor Intrusion Screening Level (VISL) calculator indicating potential risk to human health and environment.

Project Objectives

The overarching objective of this HRSC was to advance the development of an LNAPL Conceptual Site Model and to support the business goals of monetizing and controlling the long-term expense of LNAPL management for this Site.

To fulfill this business goal, the following technical objectives were established:

- Delineate and characterize, in highresolution, the lateral and vertical extent of the remaining petroleum LNAPL;
- Identify higher permeability
 heterogeneities in the subsurface that
 could serve as preferential pathways for
 LNAPL or dissolved contaminant
 migration;
- Evaluate the screened intervals of installed groundwater monitoring wells in relation to LNAPL distribution and their potential for effective recovery and monitoring of the LNAPL.
- Measure LNAPL saturation and evaluate the potential for fluid recoverability.
- Perform a preliminary screening for any potential petroleum vapor intrusion
- Identify any LNAPL present at the site boundaries

The information from this high-resolution site characterization employs "scale-appropriate" survey density to define LNAPL distribution and identify migration pathways, thereby enabling evaluation of potential risks and remedial alternatives with greater certainty. This, in turn, provides the basis to optimize management strategies and monetize the long-term management of potential risks resulting from LNAPL at this Site.

Hydrogeologic Setting

Representative boring logs drilled to install monitoring wells at the Site describe the subsurface geology consists primarily of interbedded clay and silt, with traces of sand. A layer of orange silty sand appears 12-20 feet bgs. It appears that the sand content tends to increase towards the east as drilling logs from MW-3, MW-4, MW-5, MW-6 and MW-7 show a more significant orange silty sand and sand unit encountered approximately 12-20 feet bgs.

Distinctive interbedding has created the conditions for preferential deposition and retention of LNAPL as well as preferential pathways for fluid permeability.

Groundwater levels have been measured between approximately four (4) to fourteen (14) feet below ground surface (bgs) during the period January 2017 to January 2018.

Methods, Assumptions, and Procedures

This High-Resolution Site Characterization (HRSC) was conducted in accordance with the guidelines of the Interstate Training and Regulatory Council (ITRC), Evaluating LNAPL Remedial Technologies for Achieving Project Goals (Dec 2009).

Planning for this High-resolution Site Characterization (HRSC) involved a review of available site documentation to develop an understanding of the existing Conceptual Site model (CSM) and indications of likely LNAPL impacts, and to assess the performance of remedial efforts implemented to date. Locations of monitoring wells and extraction or injection wells were loaded into the *SmartData Solutions*® real-time decision support system, along with the results of recent groundwater analyses and a high-resolution assessment work plan was developed.

Direct sensing survey station locations and the locations of soil borings advanced for high-resolution soil sampling are referenced to unique station names that identify common survey or sampling locations. In this manner, collocated LIF/UVOST® borings, MiHpt survey locations, and high-resolution soil sampling locations are referenced to the same location ID that defines the common data location. For example, the station ID "L01" will be used for both LIF direct sensing data as well as high-resolution soil samples taken at that station.

Soil sampling stations were identified to the laboratory as SB01 through 05. These have been correlated to site survey stations as:

SB01 = SF14

SB02 = SF16

SB03 = SF18

SB04 = SF07

SB05 = SF11

Laser Induced Fluorescence (LIF-UVOST®)

Utilizing Laser Induced Fluorescence (LIF-UVOST®), the vertical distribution, type, and relative concentrations of LNAPL in the subsurface can be discerned at the centimeter scale. Initial LIF-UVOST® soundings were planned to be advanced in proximity to selected wells with known free phase hydrocarbons or dissolved phase benzene or BTEX concentrations indicative of

LNAPL to characterize the presence and depth interval of LNAPL at suspect locations. The observed response of the LIF/UVOST® system at these locations would then serve as a reference for delineation of LNAPL present at the site.

The remaining LIF-UVOST® borings were planned to be advanced at selected locations stepping out from the estimated limits of the former UST enclosures and other potential release source areas, and adjacent to other wells with recent evidence of elevated petroleum hydrocarbon (PHC) impacts.

Membrane Interface Probe-Hydraulic Profiling Tool (MiHpt)

COLUMBIA Technologies employed two primary chemical detectors on the MIP for this assessment: a Photo Ionization Detector (PID) and a Flame Ionization Detector. The PID provides sensitivity to aromatic compounds (BTEX). The FID is a general detector useful for confirmation of high concentrations of organic compounds, including those not measured by the PID. Together, the two detectors provide a reliable measurement for the presence of residual petroleum LNAPL combined with concentrated adsorbed, dissolved, and vapor phase PHCs.

MiHpt soundings were advanced adjacent to monitoring wells where elevated BTEX concentrations were measured in recent and historic groundwater analyses, and adjacent to LIF/UVOST® survey locations where elevated response was measured. MIP detector response at these locations served to characterize primary areas of impact, and provide a baseline to interpret MIP response in potential areas of migration.

Using real-time information, initial survey results were evaluated to adjust the locations and depth of additional soundings advanced to isolate the primary LNAPL source area(s) from areas of PHC migration in each direction and for estimation of LNAPL boundaries.

Concurrently, HPT measurements were made to characterize relative hydraulic permeability and to identify potential migration pathways for LNAPL or dissolved phase PHCs.

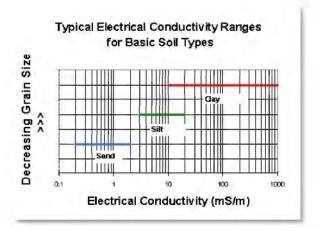
COLUMBIA Technologies employed the Hydraulic Profiling Tool (HPT) with the Electrical Conductivity (EC) system to evaluate subsurface hydrostratigraphy in the area of the release, identifying higher permeability layers or heterogeneities that constitute preferential pathways for the movement of LNAPL or dissolved-phase contaminants, and lower permeability layers that often serve as storage zones for residual hydrocarbons.

The HPT pressure logs record changes in hydraulic pressure measured directly as water is pumped into the formation at a constant rate. These logs reveal the variability and relative hydraulic conductivity of the soil.

A dipole array that measures the electrical conductivity (EC) of soil and groundwater is mounted on the tip of the MIP probe. EC measurements identify changes in the soil's electrical conductivity that can be related to changes in stratigraphy, providing insight into contaminant pathways when viewed in relation to chemical detector response.

Low EC values generally indicate coarsegrained materials (sand and gravel), while higher EC values usually indicate elevated clay content, although water chemistry and other site-specific factors influence EC response as well. General conductivity ranges for basic soil types are presented in Table 1 below (Geoprobe, 2015).

Table 1



High-resolution soil sampling

Five direct push boring locations were selected in areas where direct sensing measurements and ground water analyses indicated that LNAPL or high levels of adsorbed or dissolved phase petroleum hydrocarbons were present. Direct push borings were collocated with LIF/UVOST® and MiHpt borings at five stations SF07, 11, 14, 16, and 18.

Twenty-six (26) discrete soil samples were collected from equally spaced vertical intervals through the apparent LNAPL horizon. Target sampling intervals were selected based on adjacent direct sensing measurements. Discrete soil samples were then collected systematically from one-foot intervals from continuous core samples using TerraCore® non-methanol soil samplers in accordance with EPA Method 5035. These high-resolution soil sampling locations are also identified in **Figure 2**.

The soil samples were shipped to the ALS Environmental Laboratory to measure TPH concentrations in both the gasoline range (GRO) and diesel range (DRO) using Texas Method TX1005. Analytical reports are presented in **Appendix H**.

Results and Discussion

Hydrostratigraphy

High HPT pressure is indicative of low permeability soils. Higher permeability is manifested by low hydraulic pressure.

The HPT borings advanced at the Site show predominantly low permeability soils, consistent with the interbedded clay and silt, with traces of sand reported in borings drilled for monitoring well installation at this Site and high-resolution soil sampling conducted for this assessment.

Interbedded zones of higher permeable soils were noted in the upper fifteen (15) feet of the Site as shown in **Figure 6**.

EC measurements were also indicative of predominantly low permeability soils

LNAPL Distribution

Figure 3 presents an estimated footprint of the LNAPL at multiple depths, based on multiple lines of evidence that include LIF/UVOST® response, MIP-PID response, elevated BTEX concentrations in groundwater, and discrete confirmation soil sampling. MIP-FID response correlates with the MIP-PID confirming these data.

The vertical distribution of LNAPL is approximately six (6) to fifteen (15) feet bgs as presented in **Figures 4 through 8**

established from the multiple lines of evidence developed during this and previous investigations.

The highest measured concentrations of LNAPL determined with discrete soil sampling occurred at station SF14 just east of the diesel UST enclosure. Total Petroleum Hydrocarbons (TPH) in the C12-C28 range were measured at concentrations of 360 mg/kg and 330 mg/kg at fourteen (14) and fifteen (15) feet bgs respectively. An additional 60 mg/kg and 45 mg/kg were also measured in the C6-C12 range.

These data indicate the probable presence of diesel LNAPL at station SF14, while the remaining soil analytical results are indicative of gasoline.

Elevated MIP-PID or FID detector responses represent LNAPL, adsorbed-phase or high dissolved-phase PHCs, or a combination of these phases. Multiple instances of elevated MIP-PID and FID responses were noted between six (6) and fifteen (15) feet bgs.

The logs of direct sensing soundings made for this assessment are presented in **Appendices D**, **E**, and **F**.

High-Resolution Soil Sampling and Mobility Assesment

As discussed above, twenty-six (26) discrete soil samples were collected from equally spaced vertical intervals through the apparent LNAPL horizon at five stations SF07, 11, 14, 16, and 18 using TerraCore® non-methanol soil samplers in accordance with EPA Method 5035. These high-resolution soil sampling locations are also identified in Figure 2.

TPH analyses quantified GRO in six (6) of twenty-six (26) samples as presented in

Table 2. Soil sampling locations and depths, and GRO concentrations are depicted in

Figure 7 in comparison to the collocated LIF/UVOST® and MiHpt measurements.

Table 2 - Comparison of Measured TPH Concentrations to API Residual Saturation Screening Values for NAPL Mobility

	SOIL	C _{s at}	C _{res}	S _r
LNAPL		mg/kg	mg/kg	cm3/cm3
Gasoline	M-C Sand	143	3,387	0.02
Viiddle Distillates	M-C Sand	NR	8,000	0.04
Gasoline	M-F Sand	215	5,833	0.03
Gasoline	Silt to F-Sand	387	10,000	0.05
MPLES COLLECTED 2	3 March from Sac & I	Fox Truck Stop, Powh	atten KS	
ANALYSIS	SAMPLE ID	DEPTH (ft bgs)	RESULT (mg/kg)	
ТРН	SB01-SF14	11	ND	
TPH	SB01-SF14	12	ND	
C12 TO C28	SB01-SF14	13	60	
C6 TO C12	SB01-SF14	14	60	
C12 TO C28	SB01-SF14	14	360	
C6 TO C12	SB01-SF14	15	45	
C12 TO C28	SB01-SF14	15	330	
C6 TO C12	SB02-SF16	12	190	
C6 TO C12	SB02-SF16	14	140	
TPH	SB02-SF16	16	ND	
TPH	SB03-SF18	20	ND	
TPH	SB03-SF18	21	ND	
TPH	SB03-SF18	22	ND	
TPH	SB04-SF07	11	ND	
TPH	SB04-SF07	12	ND	
TPH	SB04-SF07	13	ND	
TPH	SB04-SF07	14	ND	
TPH	SB04-SF07	15	ND	
TPH	SB04-SF07	16	ND	
C6 TO C12	SB05-SF11	8	130	
C6 TO C12	SB05-SF11	9	110	
TPH	SB05-SF11	10	ND	
TPH	SB05-SF11	11	ND	
TPH	SB05-SF11	12	ND	
TPH	SB05-SF11	13	NO	
TPH	SB05-SF11	14	ND	
TPH	SB05-SF11	15	ND	0.00
TPH	SB05-SF11	16	ND	

On **Table 2**, above, the Total Petroleum Hydrocarbon (TPH) concentrations measured in soil samples collected from the Site are compared to published LNAPL saturation concentrations (C_{sat}), and residual LNAPL saturation concentrations (C_{res}) from API's Soil and Groundwater Bulletin No. 9 (Brost et al., 2000. *Non-Aqueous Liquid (NAPL) Mobility Limits in Soil*), a widely referenced study.

These TPH measurements validate the presence and extent of LNAPL defined at the Site through multiple lines of evidence that include high dissolved concentrations of benzene and other BTEX constituents in groundwater, LIF/UVOST® response, and MIP detector response.

Comparison of TPH concentrations measured in soil samples from the Site with residual LNAPL saturation concentrations (C_{res}) presented in **Table 2**, shows that GRO concentrations are generally less than the concentrations used as an indicator of potential LNAPL mobility in sands and silts. The trend of increasing residual saturation concentrations with decreasing grain size is noteworthy. Below C_{res}, capillary retention forces tend to limit LNAPL mobility.

TPH concentrations measured in soil samples from the Site are higher than published saturation concentrations (C_{sat}) for gasoline and middle grade distillates in sands and silts, and are indicative of residual, predominantly immobile, LNAPL. This implies that the overall LNAPL footprint is stable on a macro-scale. On a localized scale, however, LNAPL movement into and out of pore spaces (or monitoring wells) may persist, largely due to fluctuations in hydraulic conditions. That is, LNAPL may continue to exhibit micro-scale mobility within an LNAPL zone that is stable on a macro-scale.

The API screening values were developed for use in making conservative estimates of NAPL mobility, based on residual NAPL concentrations and residual NAPL saturation in unsaturated soils. The use of these values to screen for NAPL mobility presumes homogeneous soils and soil properties, which is never the case. Inherent geologic variability, macro-pores, and fractures will greatly affect the mobility and movement of NAPL. These factors must be recognized when these screening values are applied.

Concentrations above LNAPL saturation concentrations (C_{sat}) are indicative of the presence of LNAPL. Residual LNAPL saturation concentrations (C_{res}) are used as a screening limit below which LNAPL is presumed to be immobile.

Monitoring Wells and Injection Wells

Figure 8 compares the screened interval of the monitoring and recovery wells exhibiting the highest dissolved concentrations of BTEX measured in groundwater samples with the horizon of residual LNAPL developed from the multiple lines of evidence.

When compared to the distribution of soil permeability presented in Figure 9, it is apparent the screen intervals for these wells bridge across along both low permeability and higher permeability. The LNAPL appears to be stored in the interbedded silt and sandy soils at shallower depths. This condition will likely result in preferential recovery of clean groundwater from the higher permeable zone during pumping or extraction events while having minimal impact on the removal of LNAPL. Effective LNAPL fluid recovery will only occur in the discrete interval of LNAPL-to-water interface and only if the LNAPL transmissivity is sufficient to support recovery efforts.

Possible Evidence of Petroleum Vapor

This assessment included a screening for potential subsurface-vapor-to-air transport to the occupied facilities at Site. Soil concentrations for TPH indicative of residual LNAPL are present within thirty (30) feet of the occupied buildings and more shallow than fifteen (15) feet below grade. These conditions exceed the preliminary screening established by ITRC for potential petroleum vapor intrusion into the building.

Previous soil vapor measurements taken in January 2017 at a depth of six (6) feet below grade were reported by Terranext LLC. The results included two measurements of 489,000 ug/m3 at station SV-1 and 235,000 ug/m3 at station SV-2.

EPA has established a Reference Concentration (RfC) of 0.03 milligrams per cubic meter (0.03 mg/m3) for benzene based on hematological effects in humans. The RfC is an inhalation exposure concentration at or below which adverse health effects are not likely to occur. It is not a direct estimator of risk, but rather a reference point to gauge the potential for effects. At lifetime exposures increasingly greater than the reference exposure level, the potential for adverse health effects increases.¹

Quality Control

Each direct sensing instrument was operated in accordance with the manufacturer's standard operating procedures and the Standard Practice for Direct Push Technology for Volatile Contaminant Logging with the Membrane Interface Probe (MIP) ASTM STANDARD D7352 – 07.

Performance testing was performed on each system prior to and following each survey sounding. These procedures are outlined in **Appendix C.**

A QC review of the MiHpt logs for this project did not reveal any anomalies in the MiHpt system operation.

QC measures taken by the analytical laboratory ALS Environmental Laboratory are reported in Appendix H.

¹ U.S. Environmental Protection Agency. Integrated Risk Information System (IRIS) on Benzene. National Center for Environmental Assessment, Office of Research and Development, Washington, DC. 2009.

Conclusions

- This high-resolution assessment of the Sac & Fox Truck Stop delineated a residual LNAPL zone based on multiple lines of evidence that include LIF/UVOST®, MIP-PID and MIP-FID response, elevated BTEX concentrations in groundwater, and TPH (GRO and DRO) concentrations above soil saturation concentrations (C_{sat}).
- LNAPL was measured primarily in a vertical interval from six (6) to fifteen (15) feet bgs across the Site as shown in Figure 7.
- 3. TPH analyses quantified GRO in six (6) of twenty-six (26) samples as presented in **Table 2**.
- Both LIF-UVOST® and soil analytical results in the C12-C28 range at station SF14 indicate the probable presence of diesel fuel at fourteen (14) to fifteen (15) feet bgs.
- Residual phase PHCs were present at the southern boundary at stations MW-2 and SF22 at the most recently reported air-water interface.
- 6. Residual phase PHCs were present at the northern boundary at stations SF15, SF16, and SF19 above the most recently reported air-water interface. Confirmation soil samples taken at station SF16 confirmed TPH concentrations indicative of residual LNAPL at 190 mg/kg at twelve (12) feet bgs. Low concentrations of PHCs were previously reported in MW-5, however, the well has since been destroyed by site activities.
- Residual phase PHCs were present at the northeastern boundary of the facility at station SF19 above the most recently

- reported air-water interface. PHCs including 39 milligrams per liter (µg/L) of benzene were reported in MW-3, located further east of station of SF19.
- 8. TPH concentrations detected in these analyses, LIF/UVOST® and MIP response, HPT pressure logs, and the low permeability silty clay soil environment at the Site, provide multiple lines of evidence that characterize a residual, predominantly immobile, LNAPL source zone. This implies that the overall LNAPL "footprint" is stable on a macroscale, although localized LNAPL movement into and out of pore spaces (or monitoring wells) may persist.
- 9. Most monitoring wells are installed with screen intervals in soils exhibiting mostly low hydraulic permeability. The top elevation of the monitoring well screens are at times below the reported air-water interface as well as the bulk of the residual petroleum hydrocarbon mass. Both of these factors will likely result in groundwater concentrations for petroleum hydrocarbons being biased low.
- Soil concentrations for TPH indicative of residual LNAPL are present within thirty (30) feet of the occupied buildings and more shallow than fifteen (15) feet below grade. These conditions exceed the preliminary screening for potential petroleum vapor intrusion into the building. Previous soil vapor measurements taken in January 2017 at a depth of six (6) feet below grade were reported by Terranext. The results included two measurements of 489,000 μg/m3 at station SV-1 and 235,000 μg/m3 at station SV-2.

Recommendations

This report is intended to provide an improved LNAPL Conceptual Site Model (LCSM) In order to further evaluate and monetize the long-term management of LNAPL at the site. The reader is encouraged to consult the Interstate Training and Regulatory Council (ITRC). 2009. Evaluating LNAPL Remedial Technologies for Achieving Project Goals for viable technical options for LNAPL management going forward.

COLUMBIA Technologies recommends the following additional activities to further improve the LCSM and inform cleanup alternatives:

- Implement vapor mitigation adjacent to the occupied building to both reduce exposure to occupants but to also enhance natural source degradation of the residual petroleum
- Consider additional systematic highresolution soil sampling to corroborate final delineation of the LNAPL zone, measure the LNAPL mass and saturation levels, and monitor for additional parameters needed for remedial design.
- Review and refine the monitoring and recovery well network, based on the results of this HRSC.
- Measure and monitor the Natural Source Zone Depletion rate at the site to evaluate the cost effectiveness of natural depletion against the cost of most aggressive and expensive measures.
- Measure the groundwater oxidationreduction conditions and other parameters to support the evaluation of implementing and supporting aerobic degradation processes.

 Further evaluate remedial alternatives, to optimize and the effectiveness of LNAPL reduction and comprehensive risk management for this Site, following the guidelines of Interstate Training and Regulatory Council (ITRC). 2009. Evaluating LNAPL Remedial Technologies for Achieving Project Goals.

References

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APPENDIX A - Direct Sensing Equipment Description

LIF/UVOST® Equipment Description

The LIF system utilized for this investigation is the latest generation UVOST® system developed by Dakota Technologies, Inc. (DTI). The LIF-UVOST® system uses a highenergy laser to produce an ultraviolet light source for the detection of polycyclic aromatic hydrocarbons (PAHs).

The LIF-UVOST® system employs an excitation beam of light from a xenon chloride laser at 308 nanometers (nm), pulsed at 50 megahertz. Any residual phase PAHs present in the soil matrix will absorb this photon energy in the form of fluorescence.

This fluorescence is returned to the optical detection system via a second silica fiber optic line, measured, and recorded in real time across four 50nm wavelength bins centered at 350, 400, 450, and 500 nm.

Individual LIF-UVOST® logs consist of a primary graph of total fluorescence as a %RE test standard versus depth, an information box, and up to five waveform callouts. These callouts present the fluorescence intensity of each of the monitored wavelengths on the Y-axis [in microvolts (uV)]. The four peaks are due to the fluorescence at the four monitored wavelengths called channels. Each channel is assigned a color. Various non-aqueous phase liquids will have a unique waveform signature based on the relative amplitude of the four channels and/or the broadening of one or more of the channels.

The aforementioned wavelengths represent a common range of fluorescence associated with PAHs. Typically, the lighter fuels (jet fuel and gasoline) emit fluorescence at the

shorter wavelengths – 350 and 400 nm, while heavier, less distilled compounds such as bunker fuel or diesel fuel emit fluorescence at the longer wavelengths – 450 nm and 500 nm.

LIF/UVOST® screening is performed by pushing/hammering a shock protected optical cavity (SPOC) into the soil at the target rate of two centimeters per second (0.8 inches per second). As the SPOC is advanced, the total monitored fluorescence, as well as the intensity and duration of the fluorescence at each of the four monitored wavelengths, are recorded and displayed in real-time at one-second intervals as a function of depth.

LIF/UVOST® system data is presented as a percentage of the normalized % Reference Emitter (RE) performance standard. This standard consists of a blend of Non-Aqueous Phase Liquid (NAPL) and produces a consistent fluorescence response over the four wavelengths monitored by the LIF/UVOST® system. Collected data is then presented as the %RE. Using the same RE at each location and site allows normalization of data collected over several locations, sites, or screening events. The RE standard is provided by the equipment manufacturer and is the same for all LIF/UVOST® systems currently in operation.

Any fluorescence response is normally indicative of residual phase petroleum hydrocarbons, though some naturally occurring materials such as limestone will also fluoresce to a lesser and more monochromatic degree.

MIP/EC Equipment Description

The membrane interface probe with electrical conductivity (MIP/EC) probe is approximately 24 inches in length and 1.5-inches in diameter. The probe is driven into the ground

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at the nominal rate of 12 inches per minute using direct push technology (DPT) system Geoprobe or equivalent.

Geoprobe Systems developed the MIP/EC probe® that contains two separate systems: the soil EC tool and the MIP. EC, MIP chemical response, MIP operating parameters, the rate of push speed and temperature are collected by the MIP/EC Field Instrument and displayed continuously in real-time during each push of the probe.

EC: Soil electrical conductivity, the inverse of soil resistivity, is measured using a dipole arrangement. In this process, an alternating electrical current is transmitted through the soil from the center, isolated pin of the probe. This current is then passed back to the probe body. The voltage response of the imposed current to the soil is measured across these same two points. Conductivity is measured in Siemens/meter, and due to the low conductivity of earth materials, the EC probe uses millisiemens/meter (mS/m). The probe is reasonably accurate in the range of 5 to 400 mS/m.

The electrical properties of soil vary by geological setting. Therefore, conductivity measurements will vary both in magnitude and the relative change from one soil type to another in each geological setting. In general, at a given location, lower conductivity values are characteristic of larger particles such as cobbles and sands, while higher conductivities are characteristic of finer sized particles such as finer sand, silts, and clays. Observed conductivities significantly higher than 400 mS/m are indicative of ionic materials other than soil. Examples include saltwater intrusion, the presence of ionic chemicals from storage or injection, or potentially soil mixtures with metallic compounds.

MIP: The MIP portion of the probe is used to create high-resolution, real-time profiles of subsurface volatile organic compounds (VOCs). The operating principle is based on heating the soil and/or water around a semipermeable polymer membrane to 121 degrees Celsius (°C), which allows VOCs to partition across this membrane. The MIP can be used in saturated or unsaturated soils, as water does not pass through the membrane. Nitrogen is used as an inert carrier gas and travels from a surface supply down a transfer tubing which sweeps across the back of the membrane and returns any captured VOCs to the installed detectors at the surface. It takes approximately 60 seconds for the nitrogen gas stream to travel through 150 feet of inert tubing and reach the detectors.

COLUMBIA Technologies utilizes three chemical detectors on the MIP: a Photo Ionization Detector (PID), a Flame Ionization Detector (FID) and a Halogen Specific Detector (XSD) mounted on a laboratory grade gas chromatograph (GC). The output signal from the detectors is captured by the MIP/EC data logging system installed on a laptop computer.

The PID detector consists of a special ultraviolet (UV) lamp mounted on a thermostatically controlled, low volume, flow-through cell. The temperature is adjustable from ambient temperature to 250 °C. The 10.6-electron volt (eV) UV lamp emits energy at a wavelength of 120 nm, which is sufficient to ionize most aromatics such as BTEX and many other molecules such as hydrogen sulfide (H₂S), hexane, and ethanol whose ionization potentials are less than 10.6 eV.

The PID also emits a response for chlorinated compounds containing double-bonded carbons (e.g. halogenated ethylenes), such as trichloroethylene (TCE) and tetrachloroethylene (PCE). Methanol

and water, which have ionization potentials greater than 10.6 eV, do not respond on the PID. Given that the PID is non-destructive, it is often run first in series with other detectors for multiple analyses from a single injection.

The FID utilizes a hydrogen flame to combust compounds in the carrier gas. The FID responds linearly over several orders of magnitude, and the response is very stable from day to day. This detector responds to any molecule with a carbon-hydrogen bond, but poorly to compounds such as H₂S, carbon tetrachloride, or ammonia. The carrier gas effluent from the GC column is mixed with hydrogen and burned. This combustion ionizes the analyte molecules. A collector electrode attracts the negative ions to the electrometer amplifier, producing an analog signal, which is directed to the data system input.

The Halogen Specific Detector (XSD™) was developed to address the need for a sensitive and selective detector for halogenated compounds. The XSD is sensitive to halogen atoms including bromine, chlorine, and fluorine. This detector provides high halogen selectivity, making it an effective tool for identification and measurement of halogenated compounds in environments where other contaminants are present, such as high concentrations of hydrocarbons. The XSD is used to measure concentrations of a broad range of chlorinated volatile organics and other halogenated compounds.

The XSD detector consists of a ceramic probe, platinum wire (anode) and platinum bead (cathode) mounted inside a high-temperature reactor. The detector reactor combusts the incoming sample into a stream of air and converts halogenated organics into free halogen atoms. The free halogen atoms will then react with alkali atoms on the surface of the electrically charged platinum

bead, which functions as an electron emitter. When this reaction takes place, the current is measured and transmitted to the data system."

Unlike other halogen selective detectors, the XSD contains no radioactive sources and does not use organic solvents.

HPT Equipment Description

The HPT probe is approximately 24 inches in length and 1.5-inches in diameter. The probe is driven into the ground at the nominal rate of 12 inches per minute using a DPT rig.

The HPT probe was developed by Geoprobe Systems[®] and contains two separate systems: soil EC and the HPT. EC, HPT parameters, and temperature are collected by the HPT Field Instrument and displayed continuously in real-time during each push of the probe.

EC: Soil electrical conductivity, the inverse of soil resistivity, is measured using a Werner array arrangement. In this process, an electrical current is transmitted through the soil from two electrodes on the probe body. This current is then passed back to the probe, and the voltage response of the imposed current to the soil is measured across these points. Conductivity is measured in Siemens/meter, and due to the low conductivity of earth materials, the EC probe uses mS/m. The probe is reasonably accurate in the range of 5 to 400 mS/m.

The electrical properties of soil vary by geological setting. Therefore, conductivity measurements will vary both in magnitude and the relative change from one soil type to another in each geological setting. In general, at a given location, lower conductivity values are characteristic of larger

particles such as cobbles and sands, while higher conductivities are characteristic of finer sized particles such as finer sand, silts, and clays. Observed conductivities significantly higher than 400 mS/m are indicative of ionic materials other than soil. Examples include saltwater intrusion, the presence of ionic chemicals from storage or injection, or potentially soil mixtures with metallic compounds.

HPT: The HPT portion of the system is used to create high-resolution, real-time profiles of soil hydraulic properties, which can be used to infer permeability and hydraulic conductivity. The HPT system consists of a controller, a pump, a transfer line (trunkline) which is pre-strung through the DPT rods, a pressure transducer, a permeable screen, and a field computer.

HPT screening is performed simultaneously with the EC logging. As the tool is advanced, water is pumped through the trunkline and passes into the soil through the permeable screen. The flow is regulated as to be as constant as possible. The pressure required to inject the constant flow of water into the soil, known as the HPT pressure, is monitored by the pressure transducer and recorded on the field computer in pounds per square inch (psi) versus depth. The flow rate of the water into the soil formation is also measured and recorded in milliliters per minute (mL/min) versus depth.

Static pressure measurements (dissipation tests) can also be made by stopping at discrete intervals, allowing users to determine the static water level. The dissipation test provides an estimate of the static water level, based on the hydraulic head imposed on the probe at rest as compared to the pressure measured at the surface prior to starting each location push. Dissipation tests are best to run in coarse-grained materials (sands and

gravels) to assure that the local ambient hydrostatic pressure is measured quickly and accurately.

To perform a dissipation test, the HPT probe is advanced to a depth below the water table and the water flow is stopped. The pressure dissipation (reduction of pressure gradient caused by forcibly pumping water into the formation) is monitored until a stable value is observed. The dissipation usually takes the shape of a curve approaching an inflection point or stable value. The stable value is then used for the hydraulic pressure at that depth and can be used to estimate static water depth. The HPT software can also provide an estimate of K (a value used in hydrogeologic calculations) to provide an interpretation of the hydraulic permeability of the formation.

Depth in feet is measured and recorded using a precision potentiometer with a 100-inch linear range. The potentiometer is mounted on the mast of the DPT rig and a counterweight anchored to the foot of the rig. Measurements are recorded on the down stroke of the mast, as the tooling string is pushed into the ground, and is accurate within 1/10th of an inch. The reference elevation (depth) reported for each individual boring is established by setting the data logger to zero feet with the sensing window of the downhole probe aligned with the ground surface. True boring elevations can be established with the addition of survey data if provided for in the scope of work.

APPENDIX B – Interpretation of Qualitative Direct Sensing Data General LIF/UVOST® Log Interpretation

There are three primary characteristics of fluorescence that are considered when interpreting LIF/UVOST® data. These characteristics are:

- 1. Fluorescence intensity how brightly does the compound fluoresce,
- 2. Wavelength what color does the compound fluoresce at, and
- 3. Duration how long does the compound fluoresce at each monitored wavelength

Individual LIF/UVOST® logs consist of a primary graph of total fluorescence versus depth, an information box and up to five waveform "callouts". In the primary fluorescence graph, depth is plotted on the Yaxis and the combined total fluorescence intensity of the four monitored wavelengths is plotted on the X-axis. Total fluorescence intensity is presented as a percentage of the RE standard. Given that various PAHs fluoresce at differing intensities, there are several compounds that fluoresce brighter than the RE standard; therefore, the total %RE can exceed 100. Total fluorescence intensity is typically proportional to concentration and responds linearly as concentration increases.

While the magnitude of response of a LIF system may be indicative of the amount of contamination present, the system response should be considered only qualitative and not quantitative. The depth of the response is highly accurate and may be relied upon to

guide additional data gathering such as soil and/or groundwater sampling. Furthermore, the depth of the response in one boring location does provide a reliable indicator of a potential source(s) of contamination, particularly when compared to results from adjacent boring locations.

Waveform callouts are presented along with the left-hand side of the primary graph. These callouts present the fluoresce intensity of each of the monitored wavelengths on the Y-axis [in microvolts (µV)] and the duration of fluorescence of each wavelength on the Xaxis. No scale is given along the X-axis, however; it is a consistent 320 nanoseconds wide. The four peaks are due to the fluorescence at the four monitored wavelengths called channels. Each channel is assigned a color. Various NAPLs will have a unique waveform signature based on the relative amplitude of the four channels and/or the broadening of one or more of the channels. Callouts are selected by the operator and typically correspond to peaks on the primary graph.

The fill color of the response on the primary graph is based on the relative contribution of each of the four channels' area versus the total waveform area. This allows the viewer to discern different substances at different each depth interval based on the fill color.

See **Appendix H**: LIF/UVOST[®] Response to Various Saturated Products on Wet Sand for the expected wavelength signature for common compounds.

General MIP/EC Log Interpretation

Each MIP/EC log includes five separate graphs of data. The Y-axis on all graphs is depth. The first graph displays the EC,

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measured in mS/m. Small soil conductivity values are indicative of coarser grained particles, such as sands and silty sands, and larger soil conductivities are indicative of finer-grained particles, such as clays and silty clays. The next three graphs are displays of measures of chemical detector response: PID, FID, and XSD measured in µV. These graphs are a linear scale and provide a relative comparison of total detector response between boring locations. The fifth graph displays the temperature of the MIP/EC probe as it is pushed into the subsurface.

General HPT Log Interpretation

Each HPT log, presented on an individual scale, includes three separate graphs of data. The Y axis on all graphs is depth. The first graph displays HPT pressure in psi and flow rate measured in mL/min. In general, higher HPT pressure readings and lower flow rates indicate lower soil permeability, while lower HPT pressure readings and higher flow rate readings indicate higher soil permeability. The second graph shows estimated K value, in feet/day, indicating the hydraulic permeability of the formation. The static groundwater level is also displayed on the graphs. The third graph displays the EC, measured in mS/m. Lower soil conductivities are indicative of coarser grained particles, such as sands and silty sands, and higher soil conductivities are indicative of finer grained particles, such as clays and silty clays.

The HPT pressure and electrical conductivity can be used to identify hydraulic permeable layers, confining units and preferential migration pathways. This information is useful for creating contaminate fate and transport models, selecting monitoring well location and screen intervals, and targeting zones for remedial injections.

Interpreting LIF/UVOST® and Comparison to Laboratory Analyses

Generalized correlation between LIF/UVOST® and laboratory analytical results can be inferred, but cannot be viewed as a linear comparison. LIF/UVOST® response and laboratory results are collected, analyzed and reported in different units and by different procedures, so correlation is not an exact one-to-one comparison. The LIF/UVOST® uses a process whereas a 2D soil surface is exposed to excitation light, and any fluorescent light emitted is analyzed at the ground surface. Soil and groundwater results involve the collection of a soil core, extraction of sub-sample at the surface, and then transporting them to a laboratory for extraction and analysis. These processes are different by definition.

Interpreting MIP Results and Comparison to Laboratory Analyses

A typically configured MIP system is effective at profiling the relative distribution of certain VOCs and relative soil types versus depth. The typical MIP system will detect VOCs with boiling points of 121 °C or less; with vapor pressures above approximately 0.14 psi; and with non-polar hydrophobic compound structures. The sensitivity or in-situ detection level of a MIP system is dependent on many different factors. **COLUMBIA Technologies**' systems and protocols are standardized to provide reliable and comparable detection and logging of chlorinated VOCs (CVOCs) on the order of 200 ppb in-situ concentrations. Petroleum based VOCs are reliably logged at 1 ppm in-situ concentrations. Each of COLUMBIA Technologies' MIP system configurations is performance tested prior to

use and if requested, MIP systems may be specially configured for atypical compounds of concern (COCs) and site conditions.

An understanding of the principles of operation and performance of the configured MIP detectors is essential to properly interpret the MIP log results. For example, a CVOC with an ionization potential greater than 10.6 eV will respond on the XSD detector but not on the PID equipped with a 10.6 eV lamp. A hydrophilic compound such as an alcohol or ketone will normally be scrubbed out of the MIP gas stream by the MIP Membrane and the installed dryer and never reach the detectors. A CVOC with a small number of chlorine atoms such as vinyl chloride or DCE will have a lower response on the XSD than a CVOC containing three or four chlorine atoms. Properly configuring and testing the MIP system for the site-specific COCs prior to use can overcome each shortfall in detector or system performance. Additionally, the in-field performance tests performed before and after each boring are critical to monitor the performance of the MIP system from the membrane through to the data logging system.

Generalized correlations between MIP response and laboratory sample results can be inferred, but cannot be viewed as a linear comparison. MIP response and laboratory results are collected, analyzed and reported in different units and by different procedures, so correlation is not an exact one-to-one comparison. For example, not all VOCs present and analyzed in laboratory instruments with compound separation are detected and measured by a typical MIP system. The MIP process uses a membrane extraction process from a heated zone of varying subsurface matrix of soil, water, and/or vapor. Soil and groundwater results involve the collection of a sample, extraction

of sub-sample at the surface, and then transporting them to a laboratory for further extraction and analysis. These two processes are different by definition.

Unusual or invalid responses on the MIP system can result from malfunctions such as carrier or makeup gas leakage, gas flow blockage, heater failure, and carryover of water vapor or excessive chemical saturation. Each MIP detector will respond differently to each of these malfunctions. The most common cause of false positive responses for CVOCs is water carryover or blockage of carrier gas flow. The most common causes of false negative are improperly adjusted gas flows or leakage and inoperative detectors. COLUMBIA Technologies' field geochemists are trained to recognize these problems and to take the appropriate corrective action in the field.

APPENDIX C – Quality Control Procedures

LIF/UVOST® System Performance Test

As a quality control check, the LIF/UVOST® system response is evaluated prior to and upon completion of each LIF/UVOST® screening location. This evaluation is completed using a RE that consists of a blend of NAPL and produces a consistent fluorescence response over the four wavelengths monitored by the LIF/UVOST® system. Collected data is then presented as a percentage of the RE. Using the same RE at each location and site allows normalization of data collected over several locations, sites, or screening events. The RE standard is provided by DTI and is the same for all LIF/UVOST® systems currently in operation.

In addition to obtaining a baseline RE for each location, the background reading of the LIF/UVOST® system is electronically recorded prior to insertion into the soil. This background reading is required to be less than 0.5% of RE prior to the start of any testing. The background during tool advancement typically stays at or below the surface background reading – giving confidence that any increases in fluorescence are "true" readings and not fluctuations or variations in background.

MIP/EC System Performance Test

As a quality control check, the MIP system response is evaluated prior to and upon completion of each MIP location. An aqueous phase performance test is performed using specific compounds designed to evaluate the sensitivity of the particular probe, transfer line and detector suite to be used. The resulting values are

recorded and compared to predetermined values.

The EC dipole is also evaluated using a brass and stainless steel test jig, resulting in known values of 55 and 290 mS. Results must fall within 10% of the expected values; otherwise, corrective action must be performed.

HPT System Performance Test

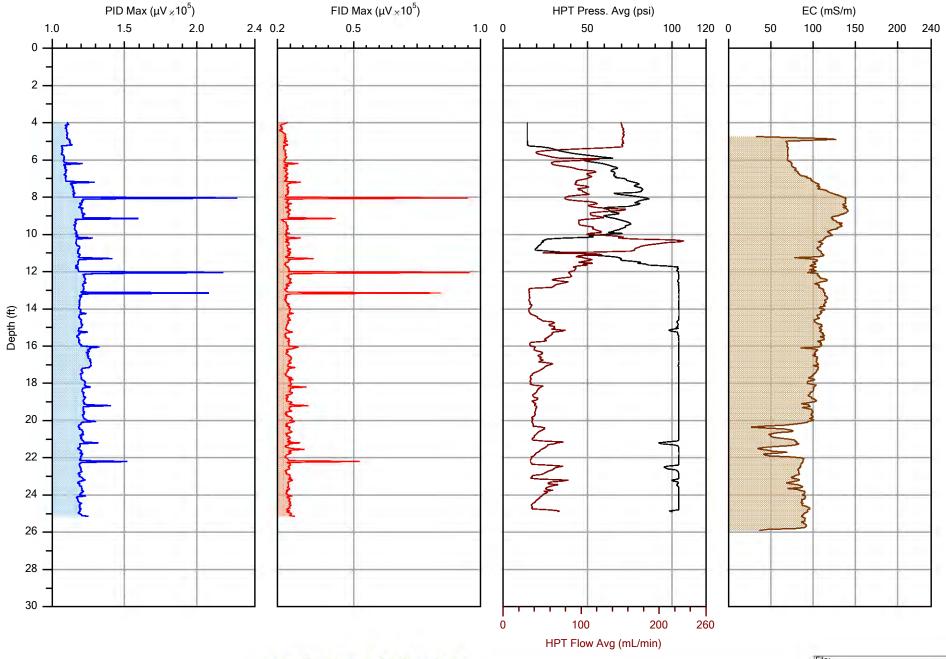
The EC Wenner is also evaluated using a Wenner Array test jig, to test the probe for isolation and continuity. Results must fall within 10% of the expected values; otherwise, corrective action must be performed.

The HPT sensor is also evaluated using static (no flow) and dynamic (with the flow at approximately 150 milliliters per minute hydraulic pressure measurements at two different head elevations, 6.0 inches apart. The difference for each test must be 0.2 psi, +/- 10%; otherwise, corrective action must be performed.

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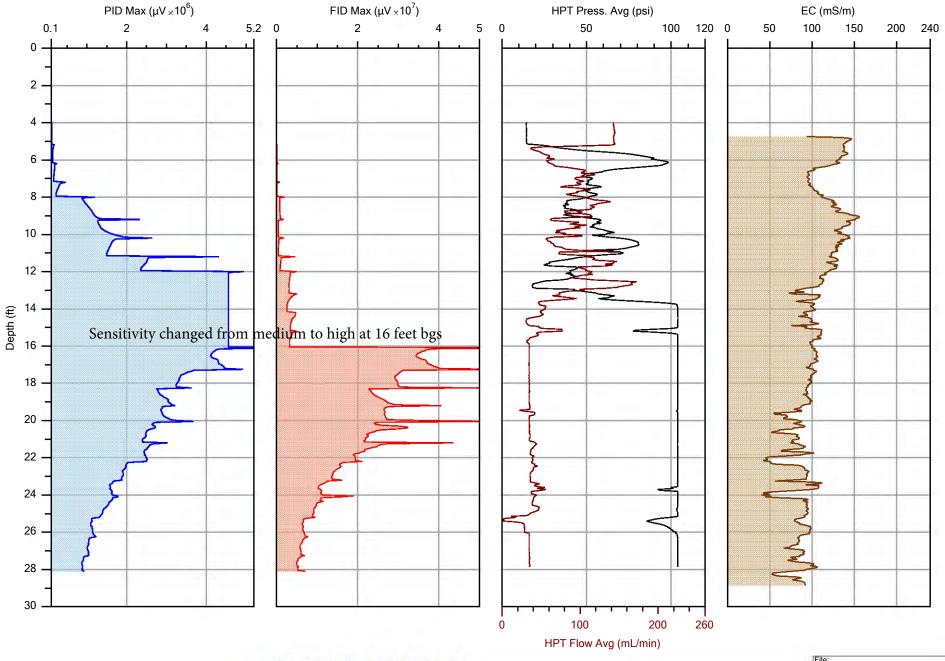
APPENDIX D – Data Logs for Membrane Interface Probe/EC with Hydraulic Profile Tool (MiHpt)

Individual Scale



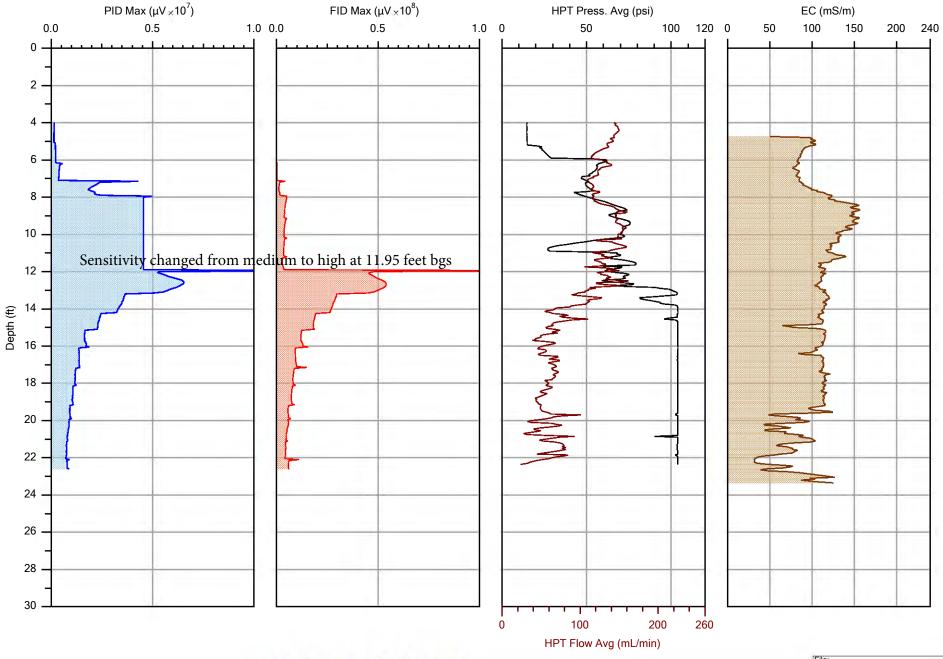


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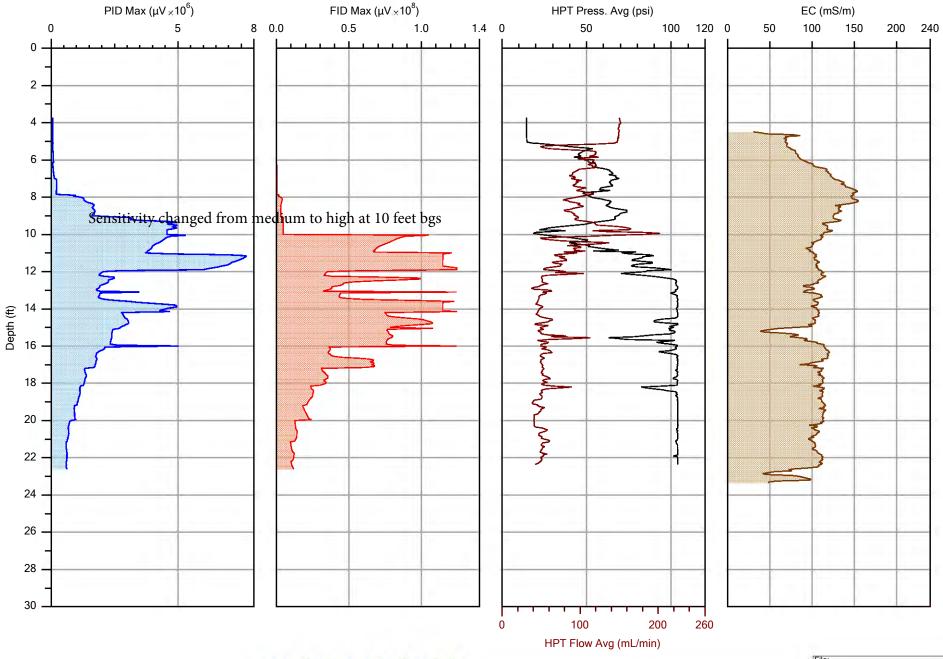


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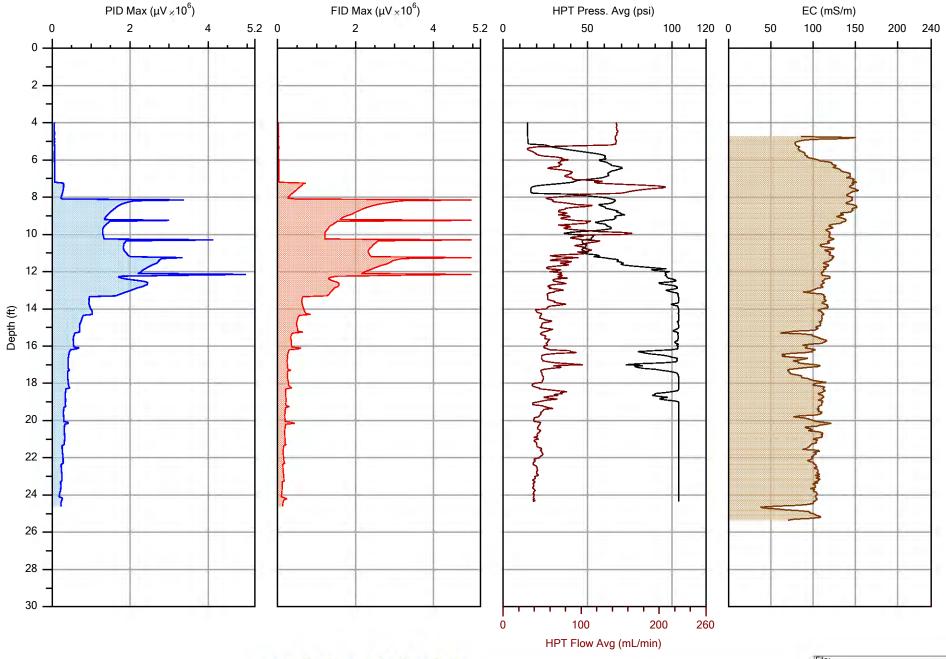


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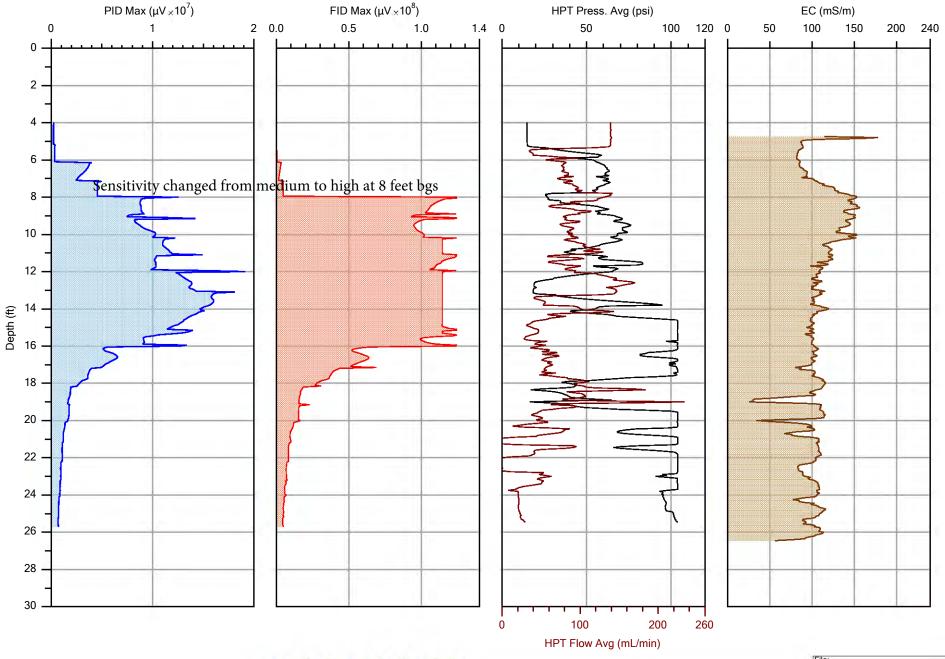


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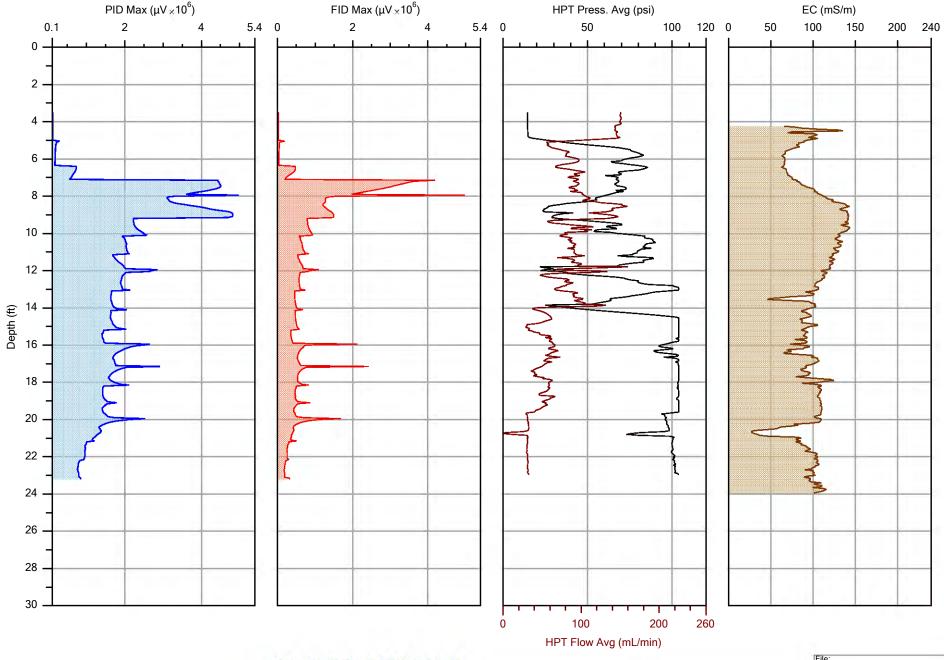


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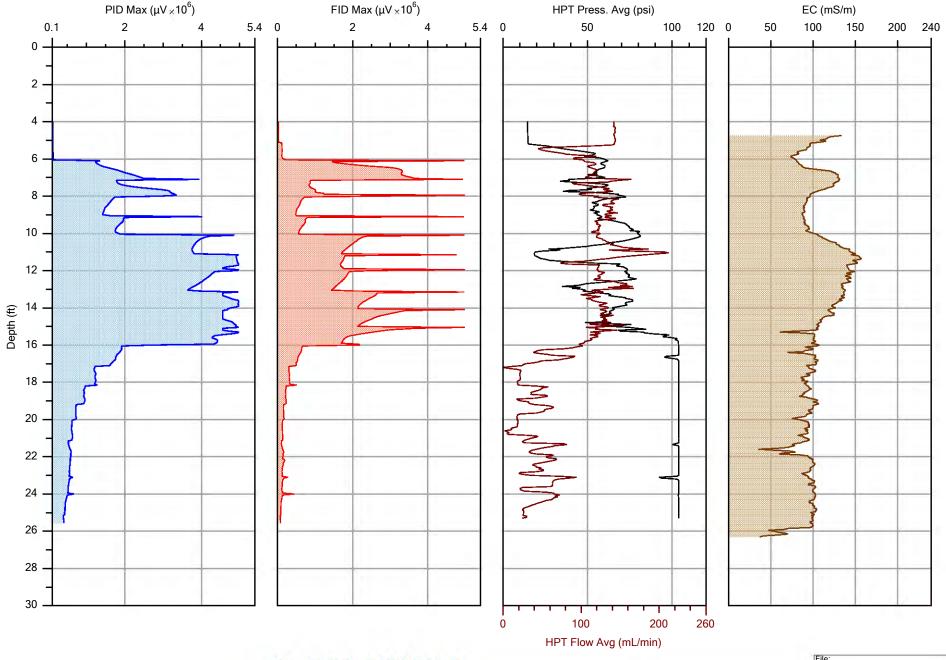


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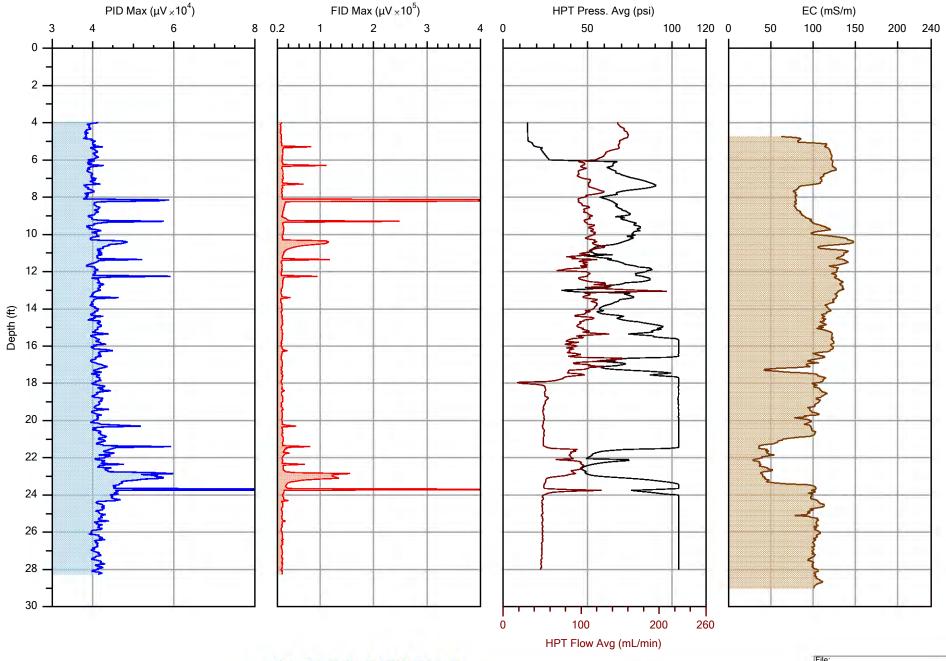


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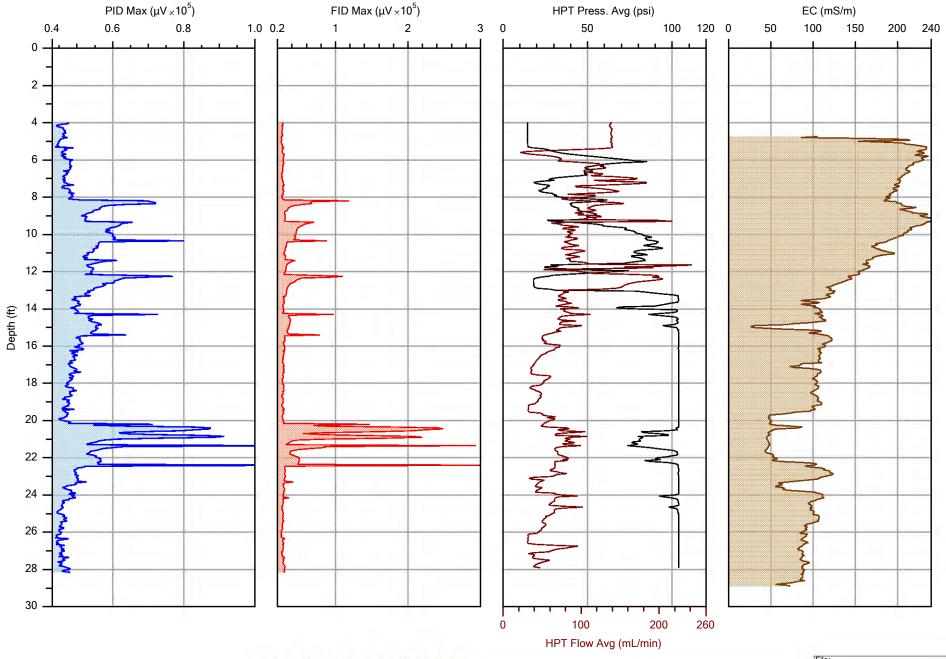


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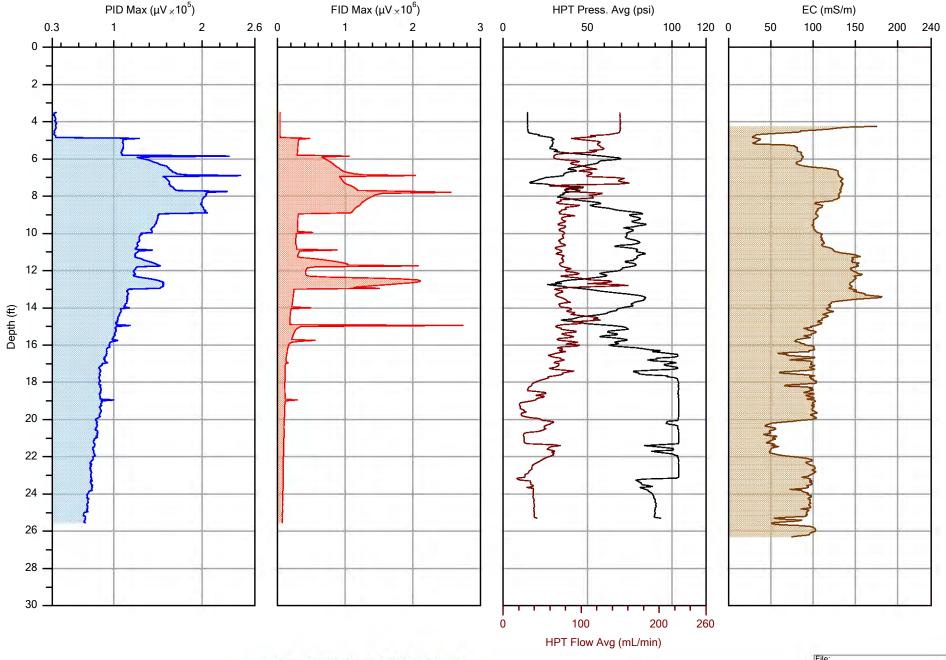


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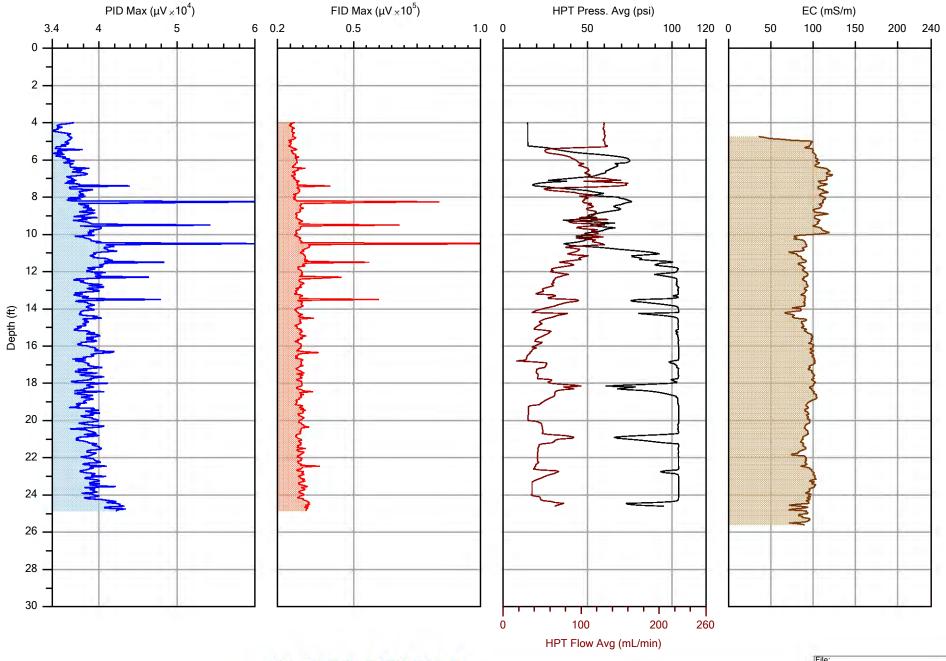


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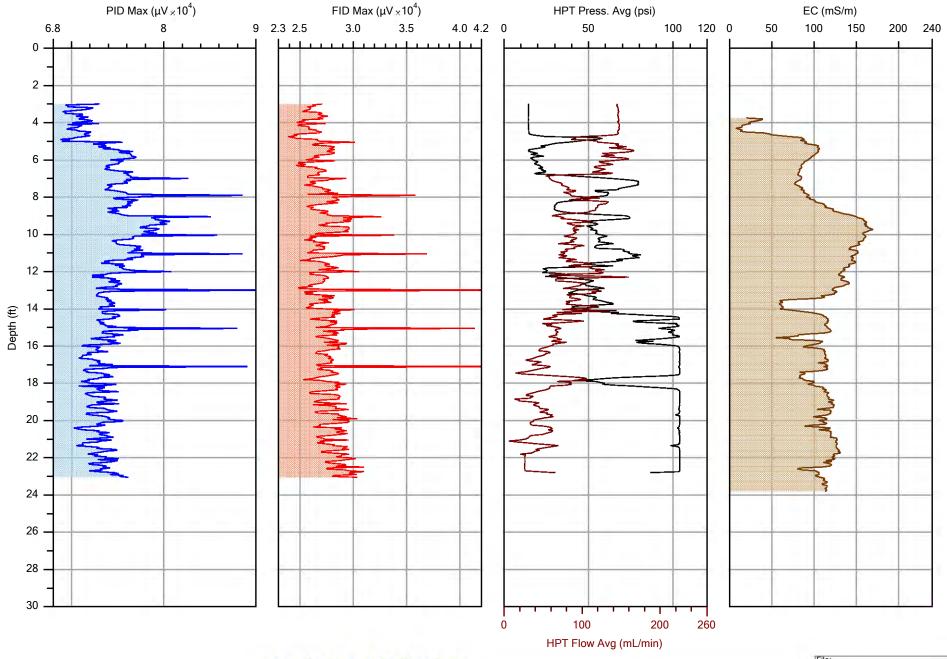


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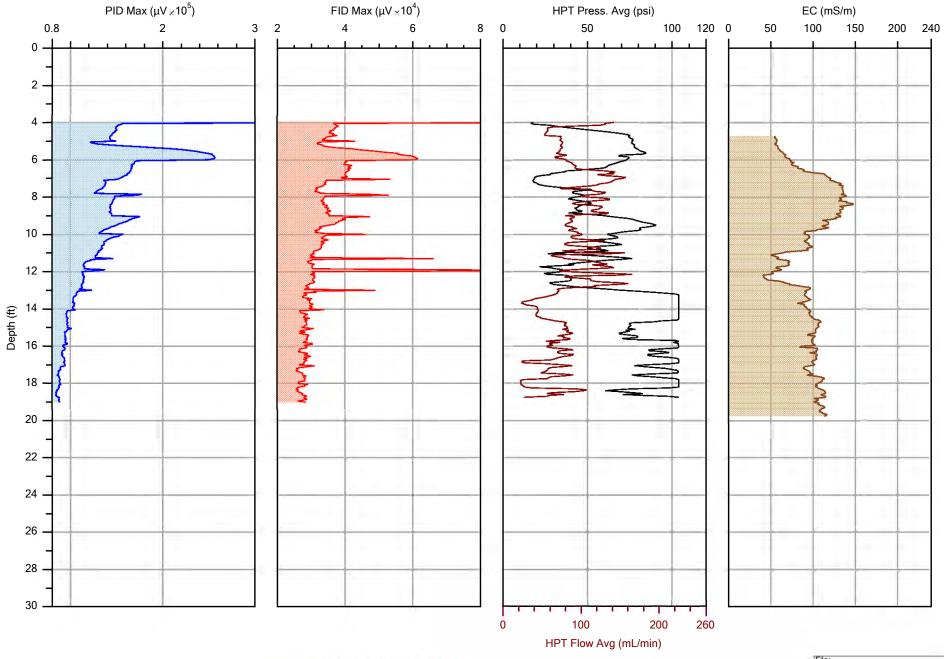


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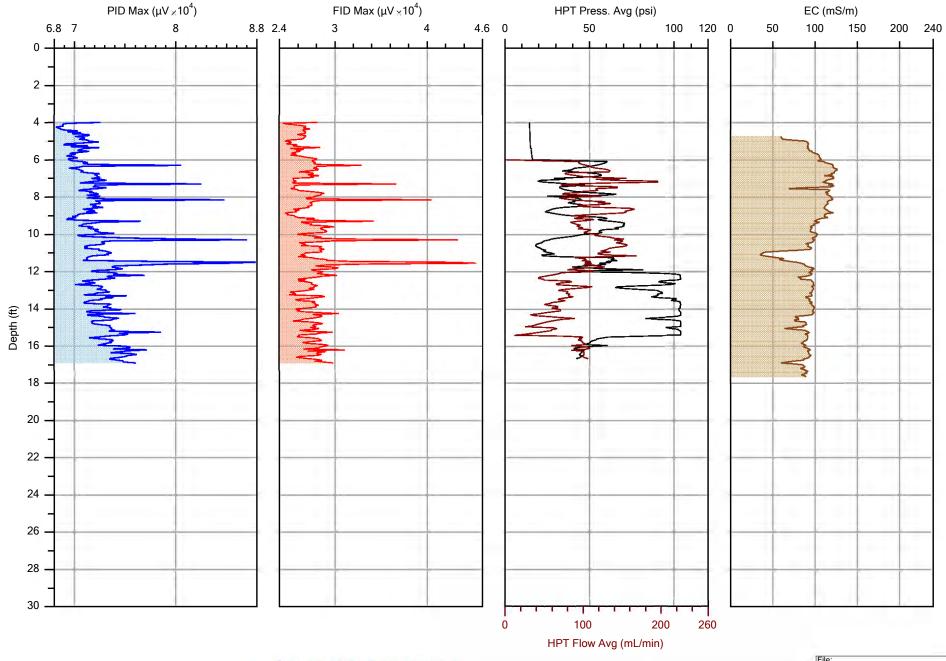


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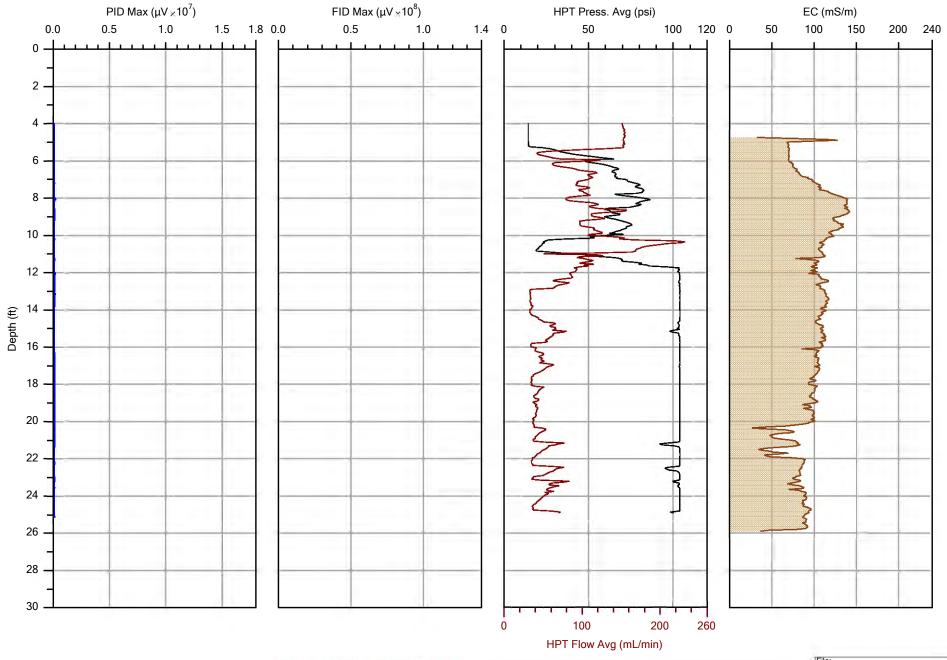




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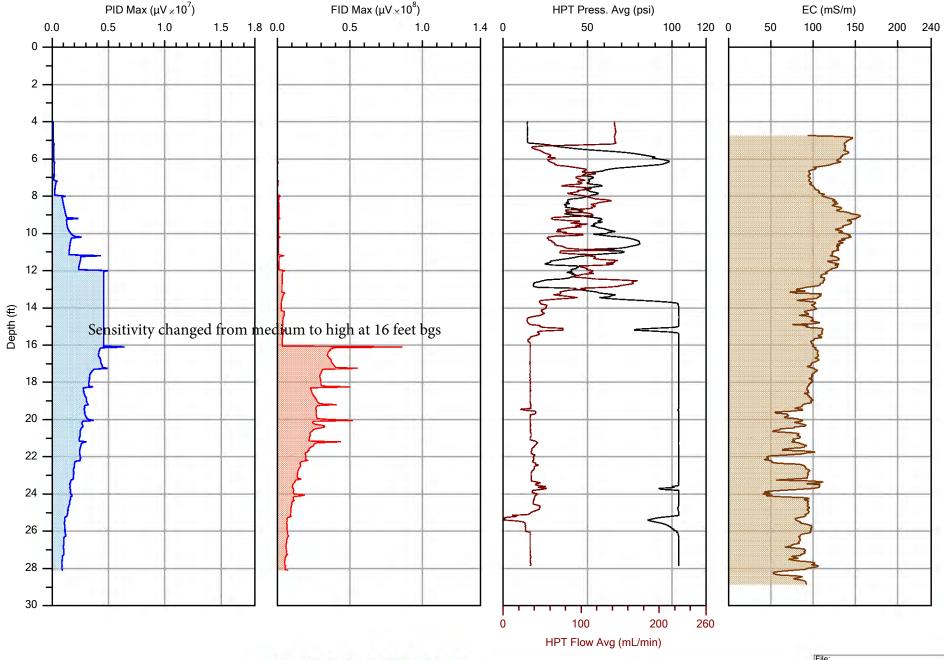
APPENDIX E – Data Logs for Membrane Interface Probe/EC with Hydraulic Profile Tool (MiHpt)

Collective Scale



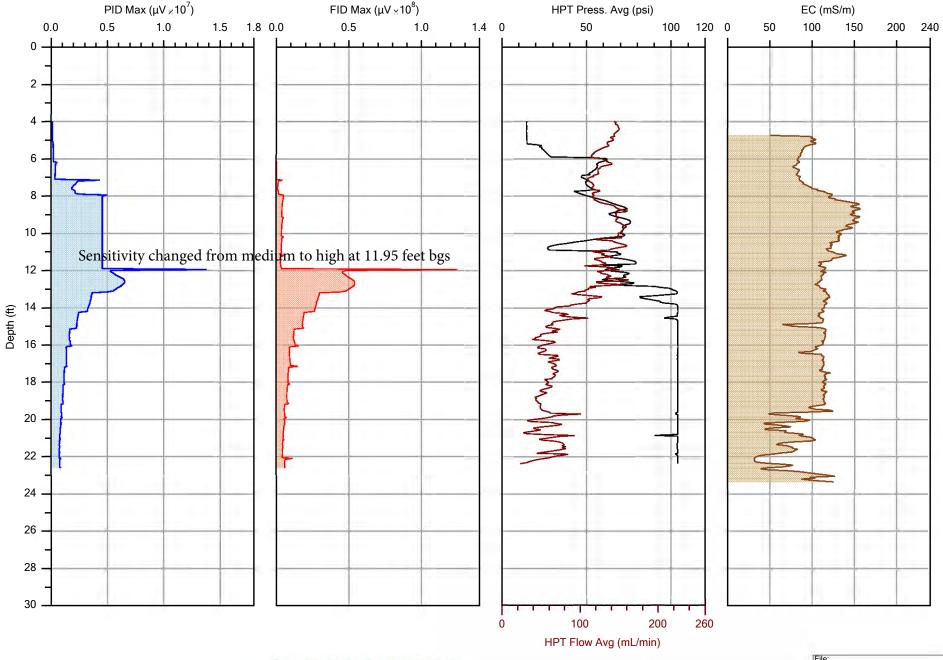


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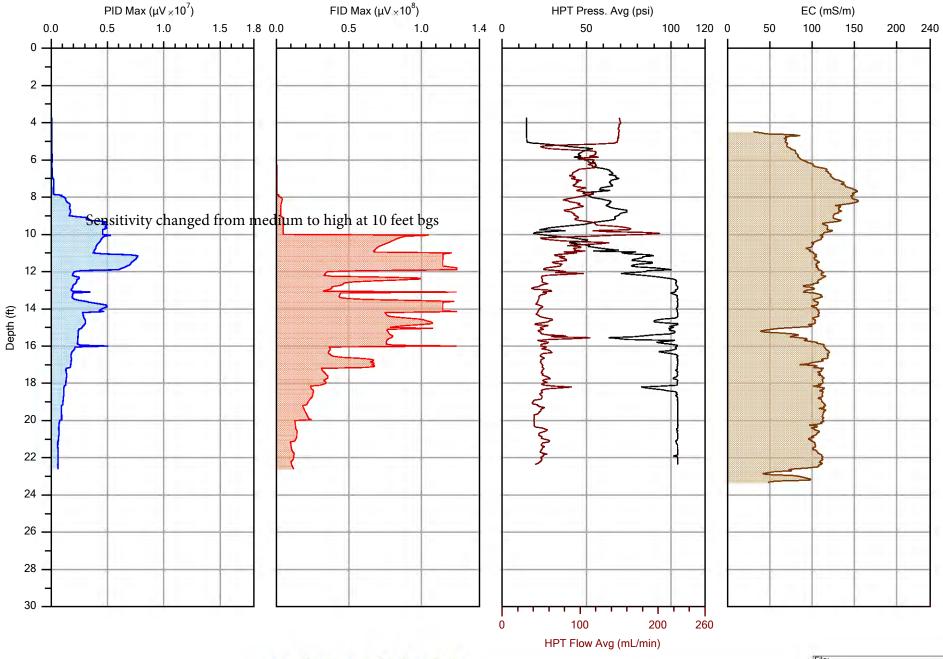


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Company:	Operator:	Date:	
COLUMBIA Technologies	AMS	3/20/2018	
Project ID:	Client:	Location:	
Sac & Fox Truck Stop	Terranext LLC	Powhattan, KS	



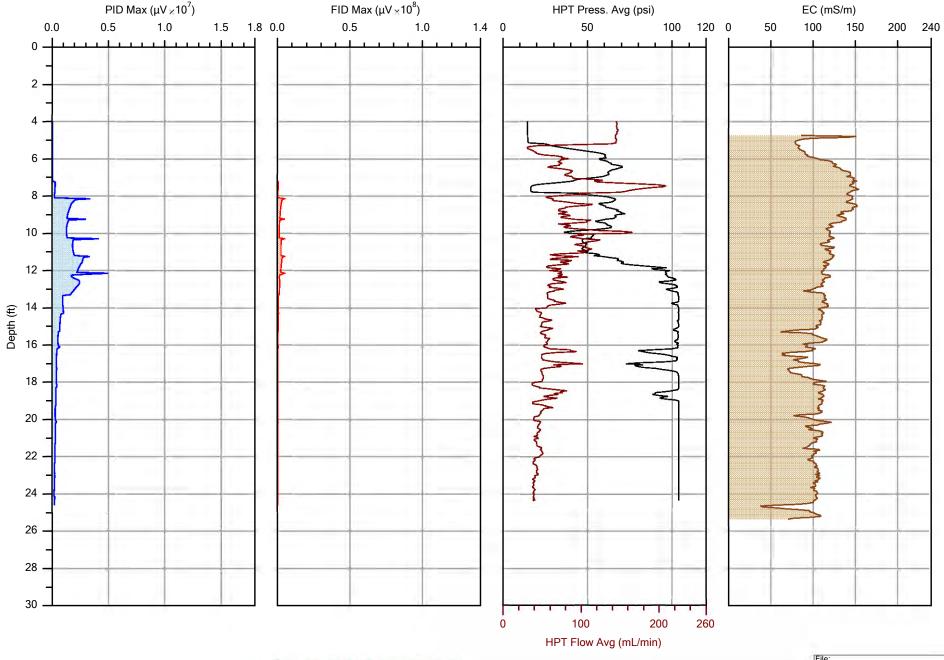


Sac & Fox Truck Stop	Terranext LLC	Powhattan, KS
Project ID:	Client:	Location:
COLUMBIA Technologies	AMS	3/20/2018
Company:	Operator:	Date:
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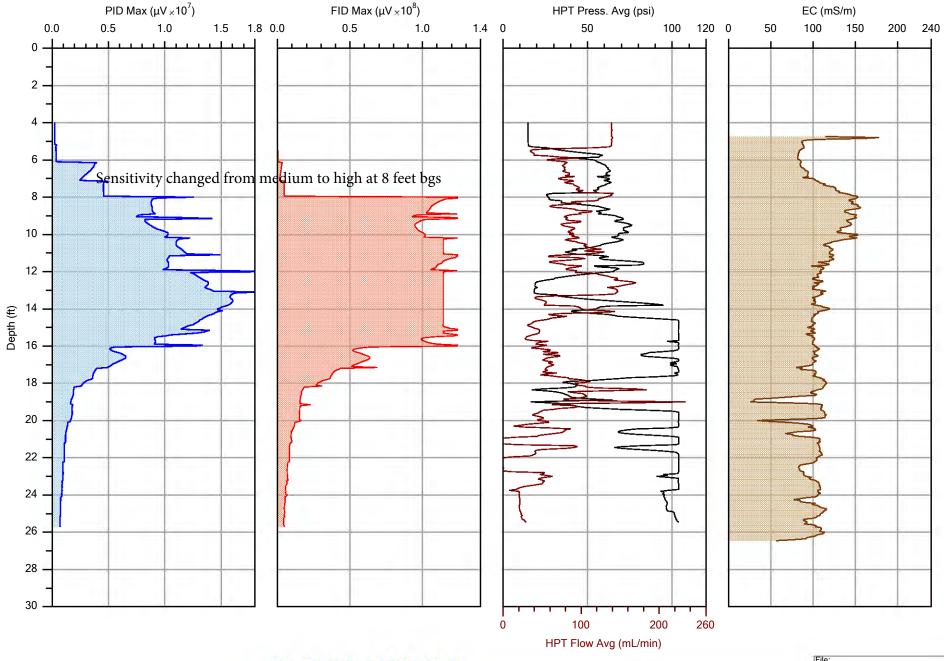


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COLUMBIA Technologies	AMS	3/21/2018
Project ID:	Client:	Location:
Sac & Fox Truck Stop	Terranext LLC	Powhattan, KS



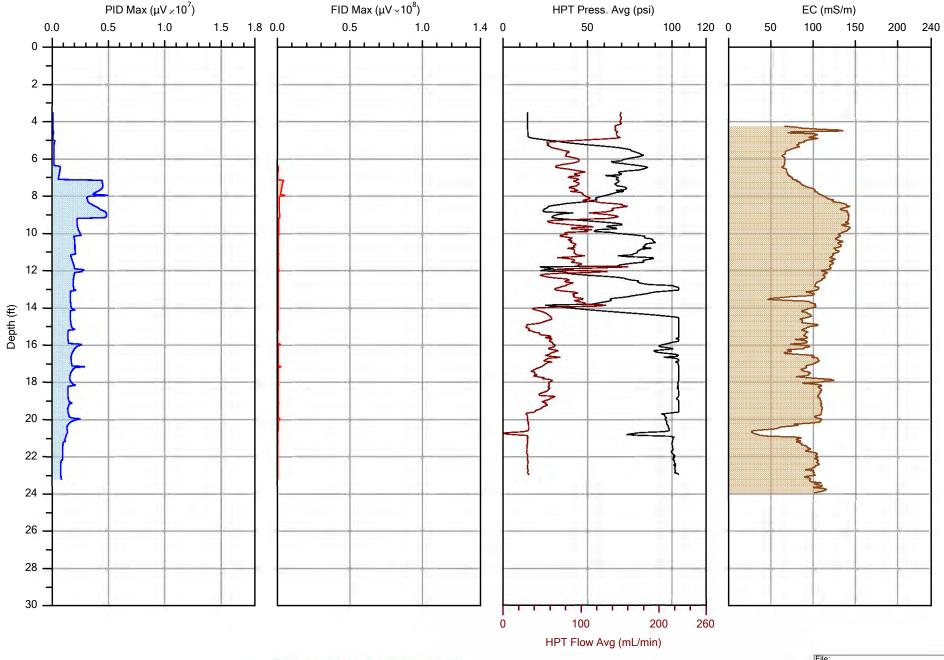


Sac & Fox Truck Stop	Terranext LLC	Powhattan, KS
Project ID:	Client:	Location:
COLUMBIA Technologies	AMS	3/21/2018
Company:	Operator:	Date:
		SF10.MHP



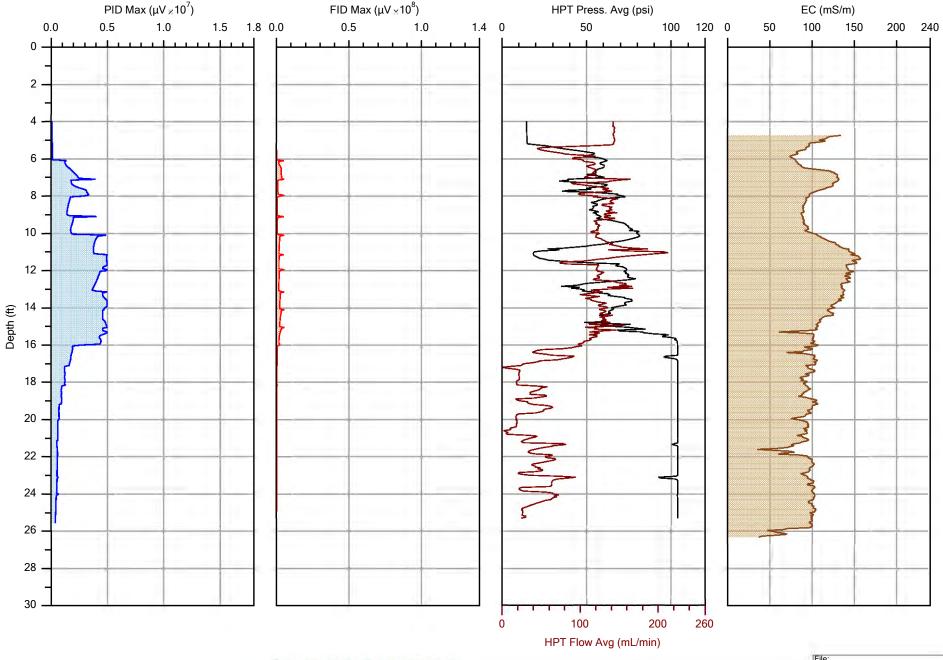


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Company:	Operator:	Date:	
COLUMBIA Technologies	AMS	3/20/2018	
Project ID:	Client:	Location:	
Sac & Fox Truck Stop	Terranext LLC	Powhattan, KS	



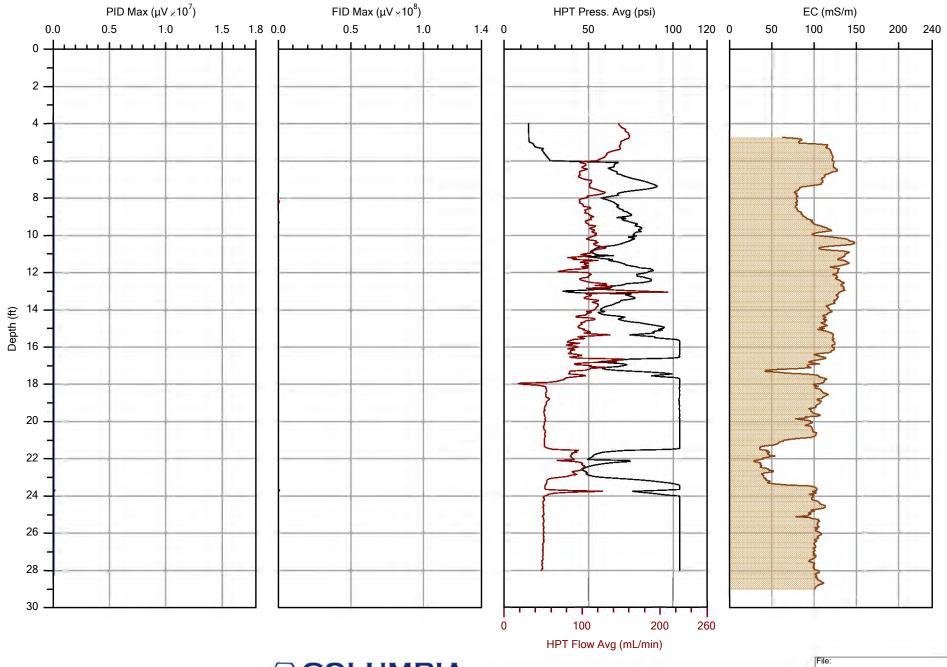


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COLUMBIA Technologies	AMS	3/20/2018
Project ID:	Client:	Location:
Sac & Fox Truck Stop	Terranext LLC	Powhattan, KS



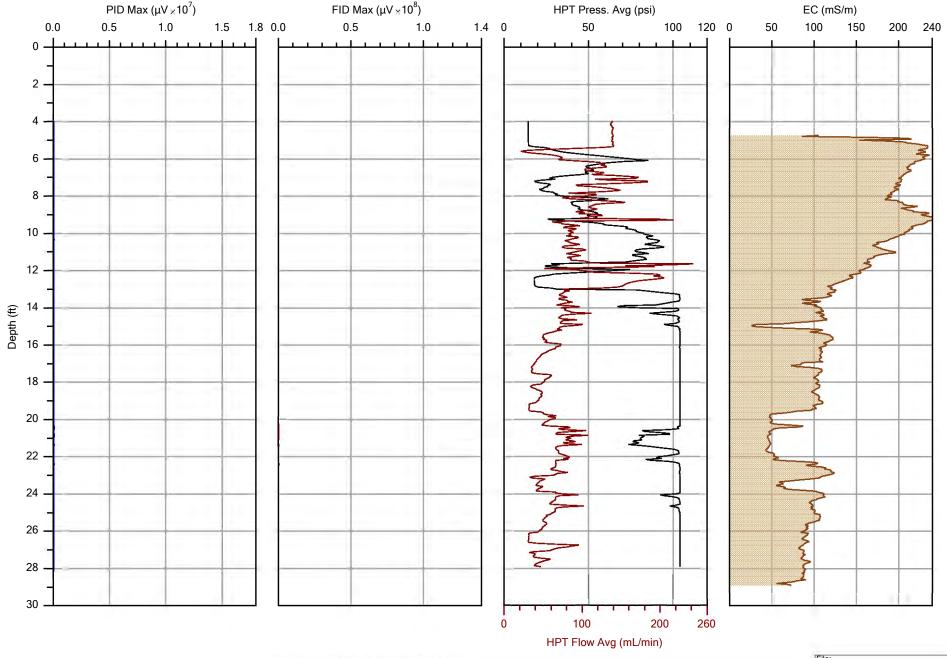


Sac & Fox Truck Stop	Terranext LLC	Powhattan, KS
Project ID:	Client:	Location:
COLUMBIA Technologies	AMS	3/20/2018
Company:	Operator:	Date:
		SF16.MHP



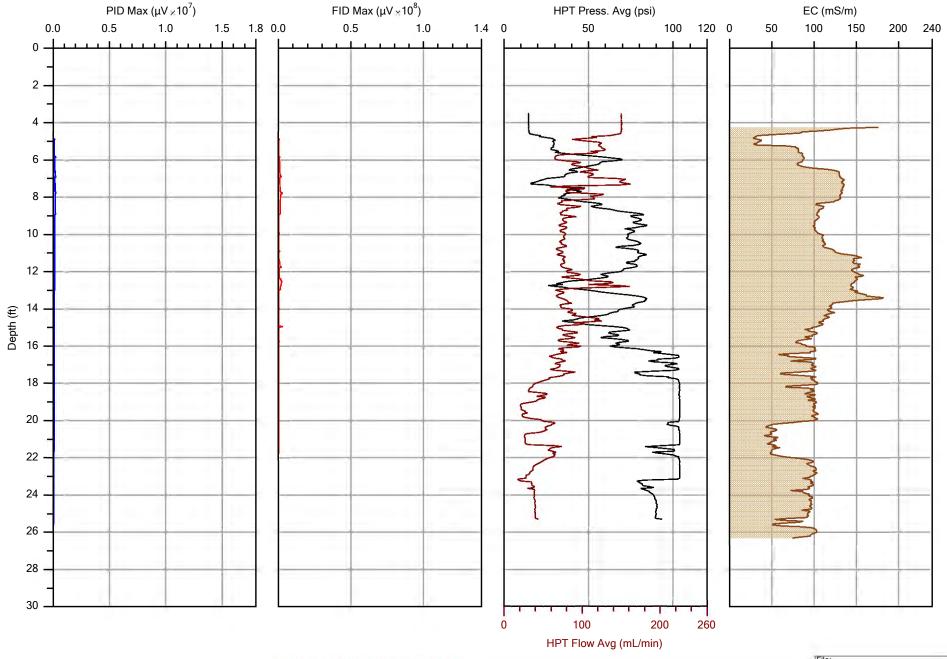


Sac & Fox Truck Stop	Terranext LLC	Powhattan, KS
Project ID:	Client:	Location:
COLUMBIA Technologies	AMS	3/21/2018
Company:	Operator:	Date:
		SF17.MHP



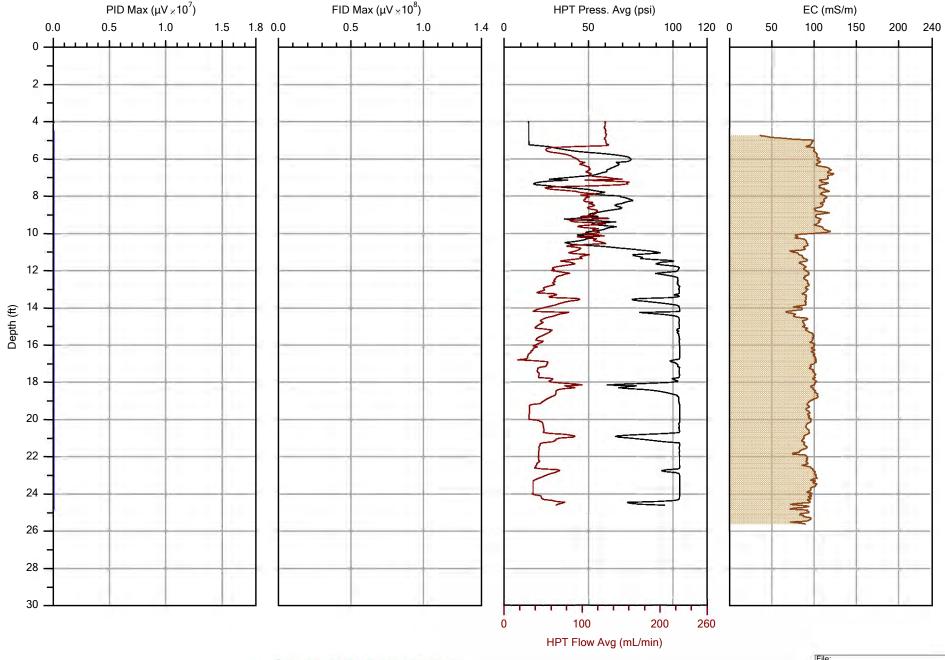


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Company:	Operator:	Date:	
COLUMBIA Technologies	AMS	3/21/2018	
Project ID:	Client:	Location:	
Sac & Fox Truck Stop	Terranext LLC	Powhattan, KS	



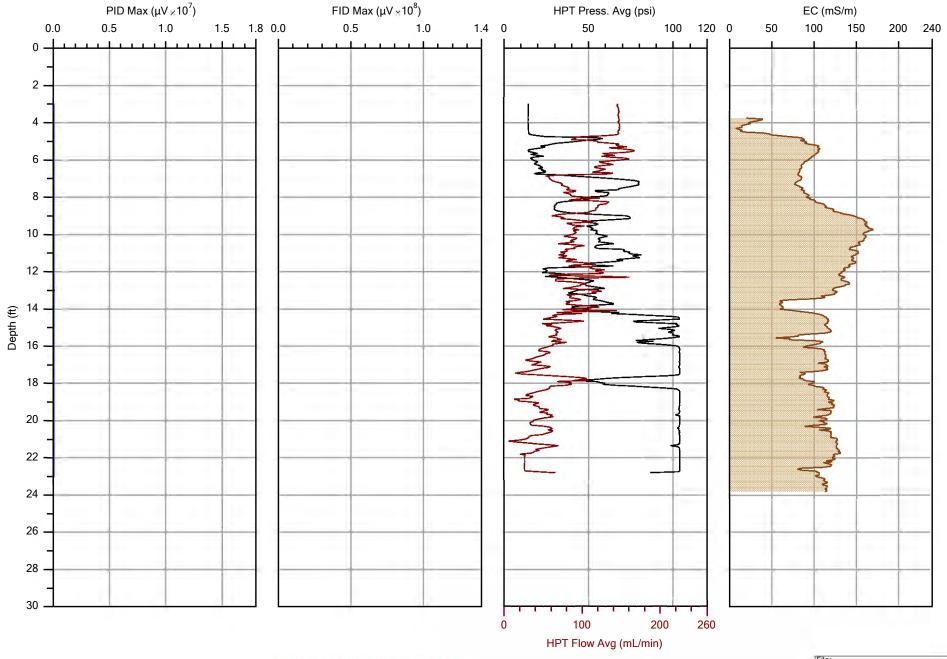


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COLUMBIA Technologies	AMS	3/21/2018
Project ID:	Client:	Location:
Sac & Fox Truck Stop	Terranext LLC	Powhattan, KS



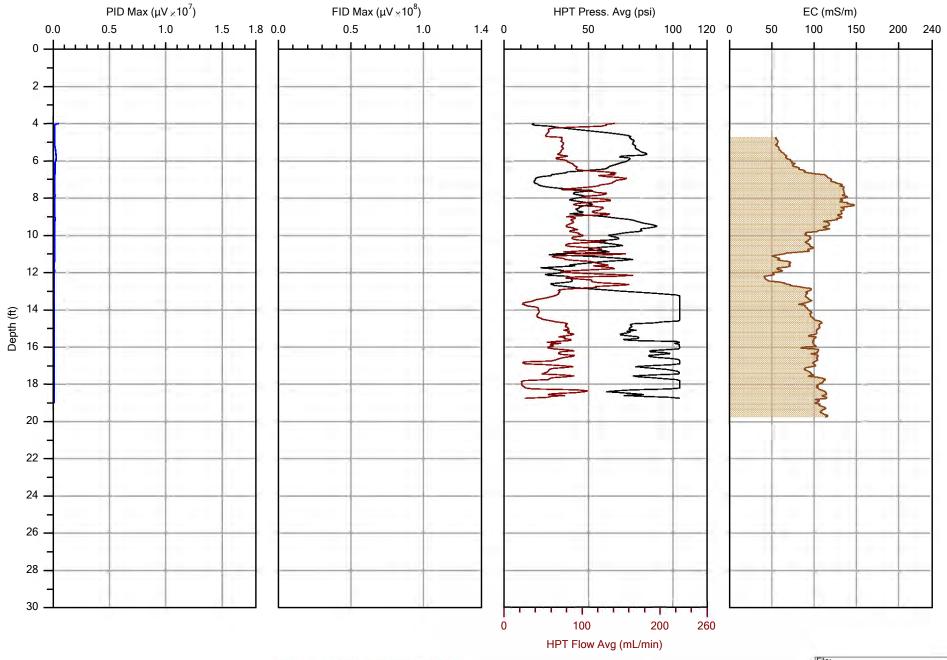


Sac & Fox Truck Stop	Terranext LLC	Powhattan, KS
Project ID:	Client:	Location:
COLUMBIA Technologies	AMS	3/21/2018
Company:	Operator:	Date:
		SF20.MHP



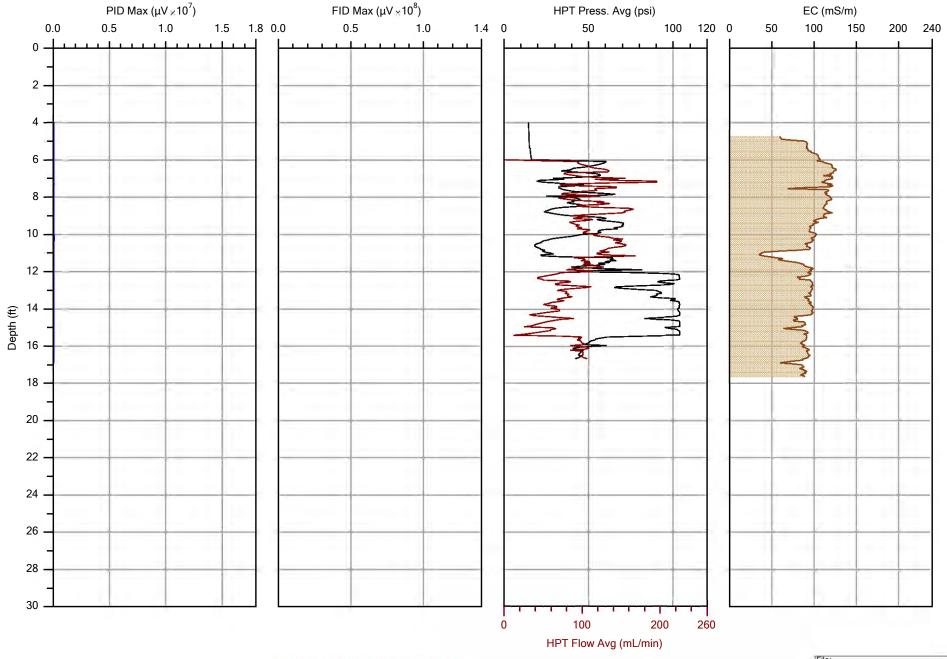


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COLUMBIA Technologies	AMS	3/22/2018	
Project ID:	Client:	Location:	
Sac & Fox Truck Stop	Terranext LLC	Powhattan, KS	





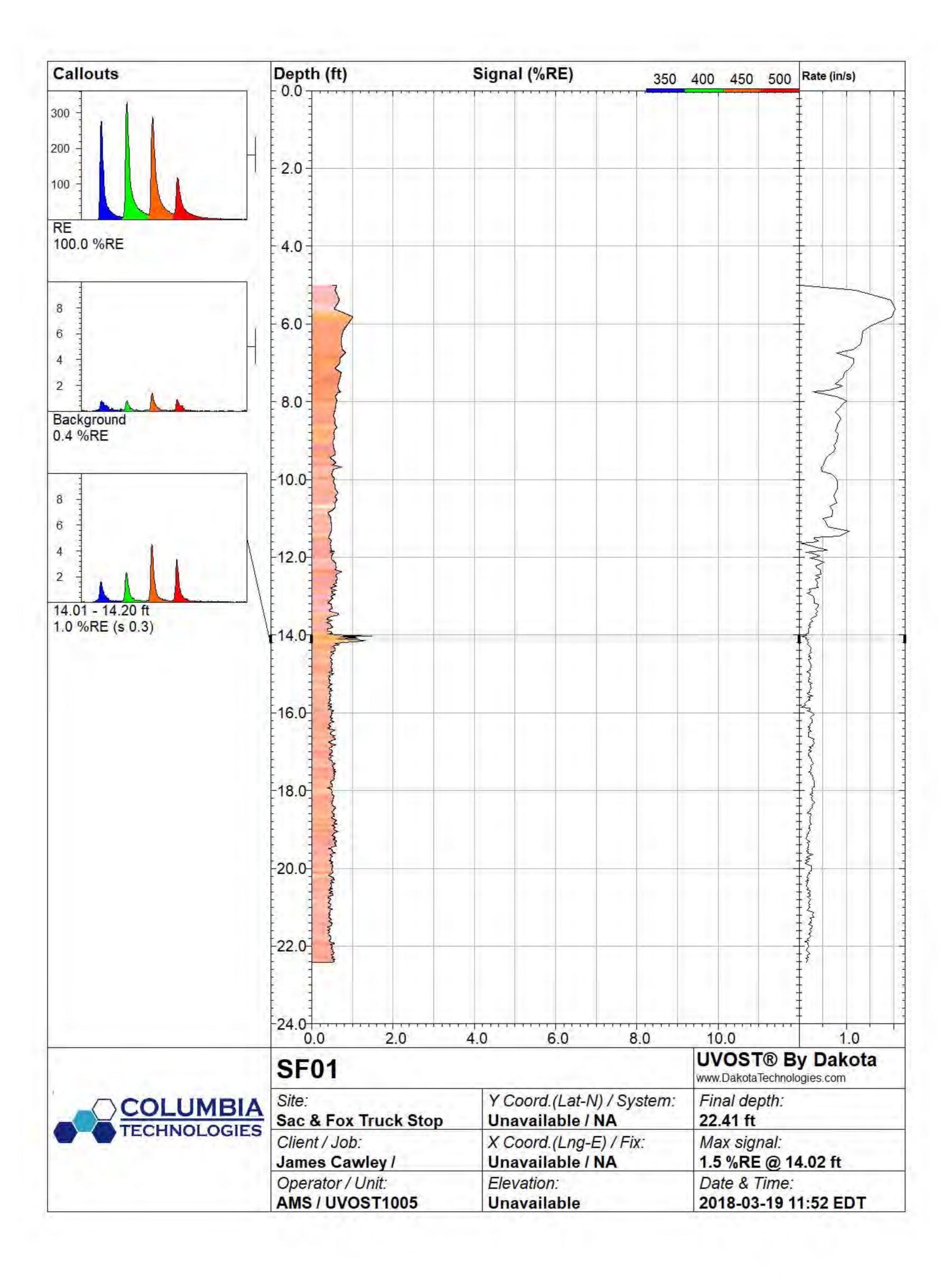
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Company:	Operator:	Date:
COLUMBIA Technologies	AMS	3/22/2018
Project ID:	Client:	Location:
Sac & Fox Truck Stop	Terranext LLC	Powhattan, KS

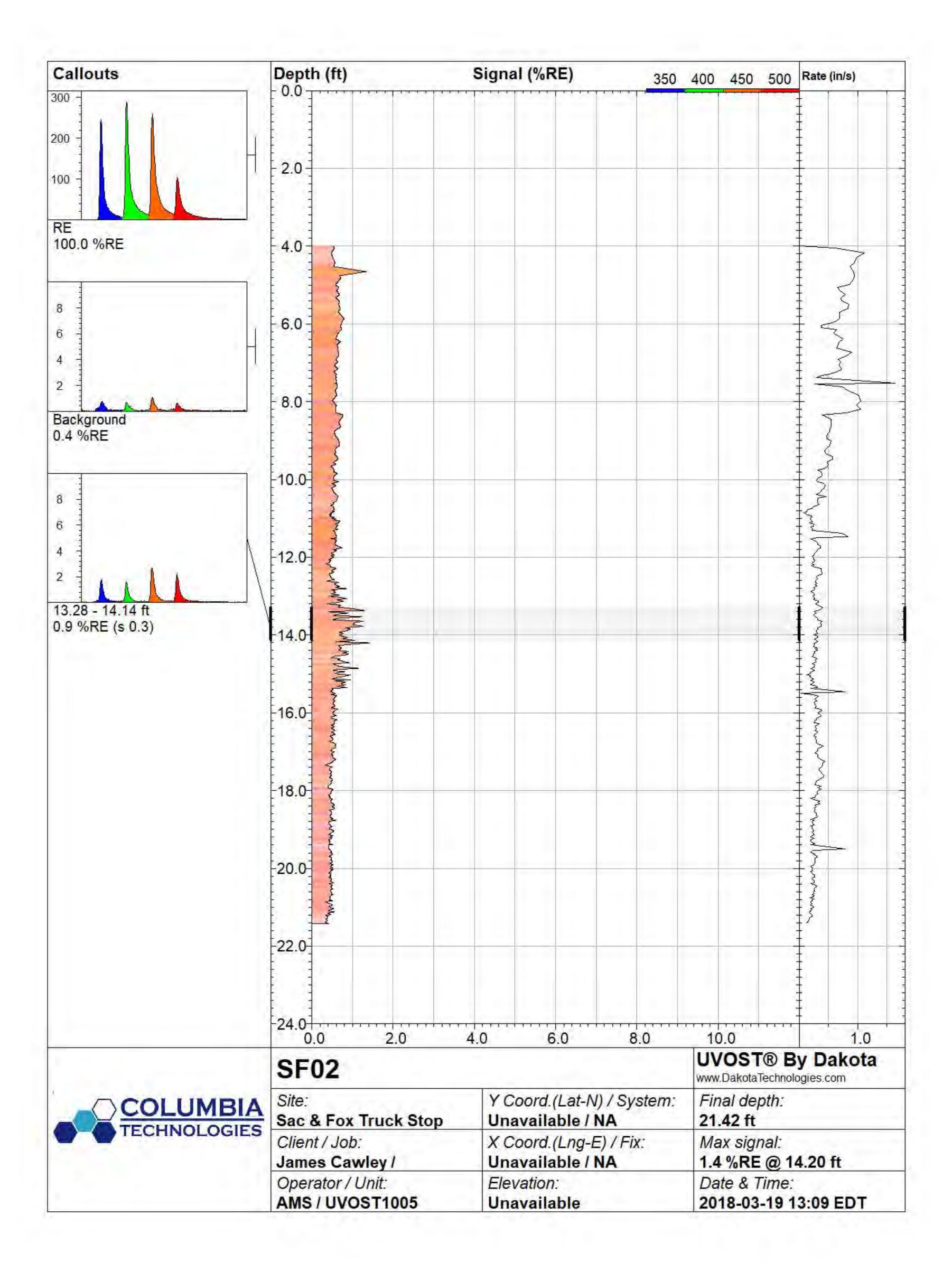


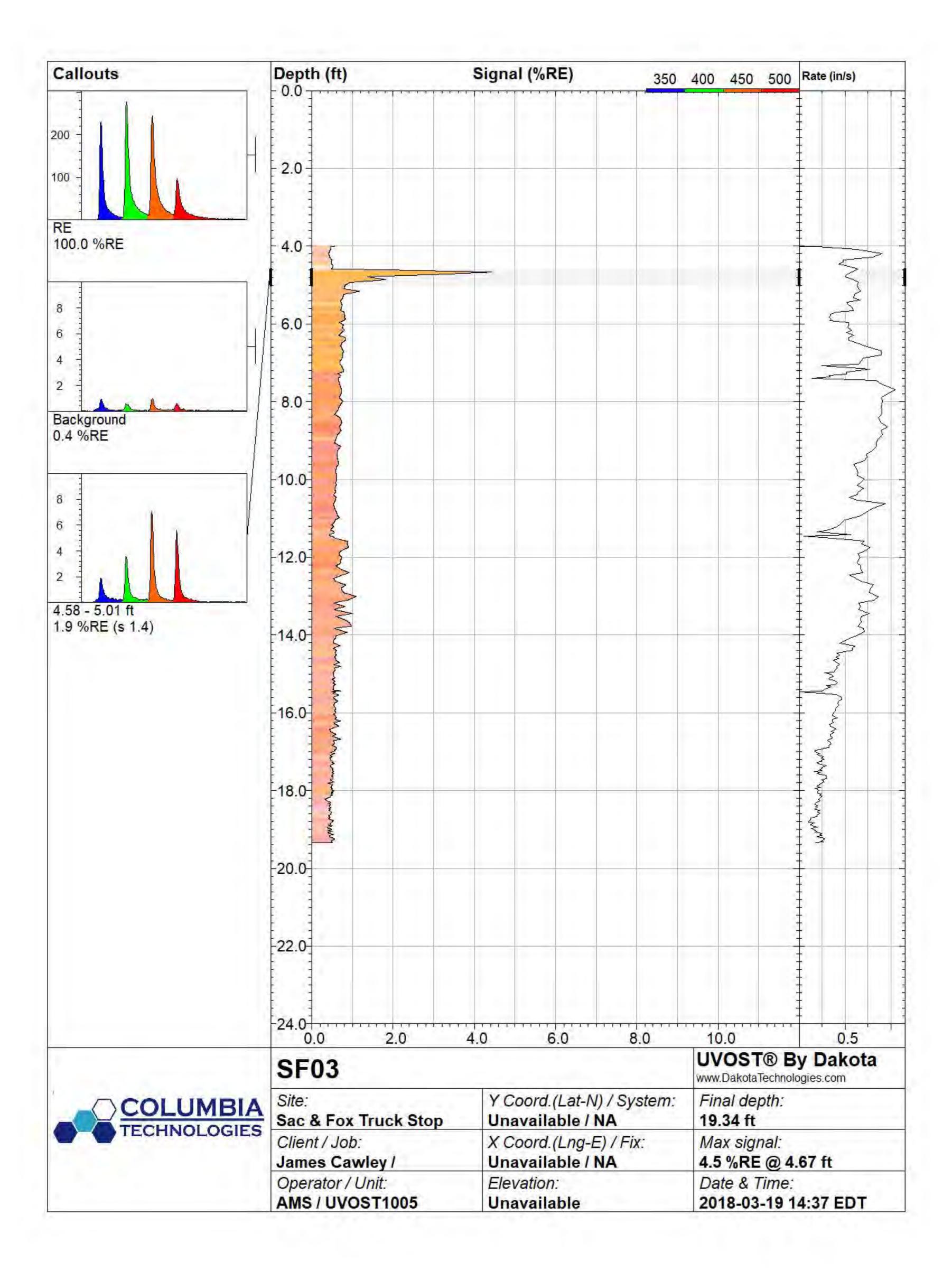


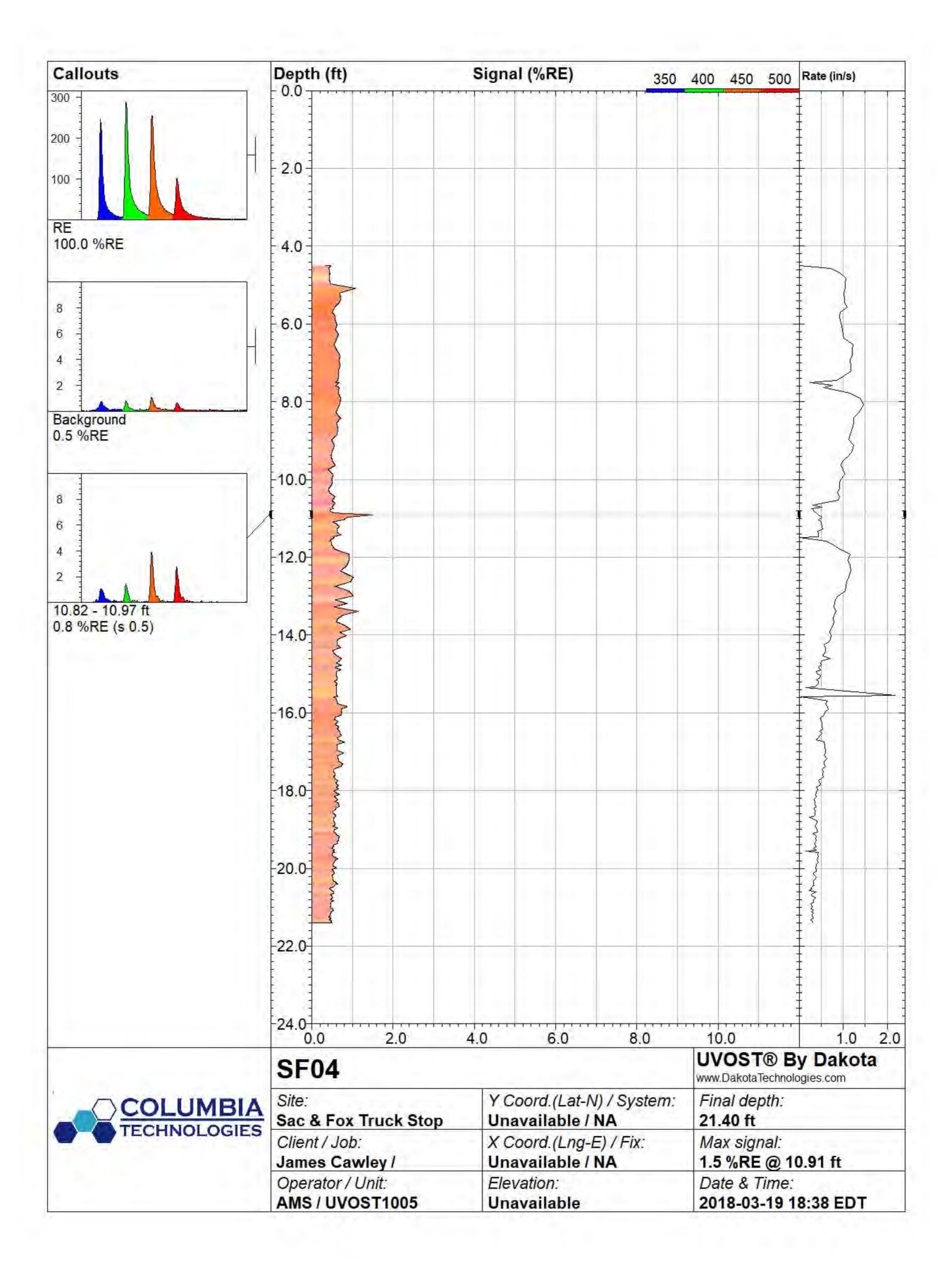
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Company:	Operator:	Date:	
COLUMBIA Technologies	AMS	3/22/2018	
Project ID:	Client:	Location:	
Sac & Fox Truck Stop	Terranext LLC	Powhattan, KS	

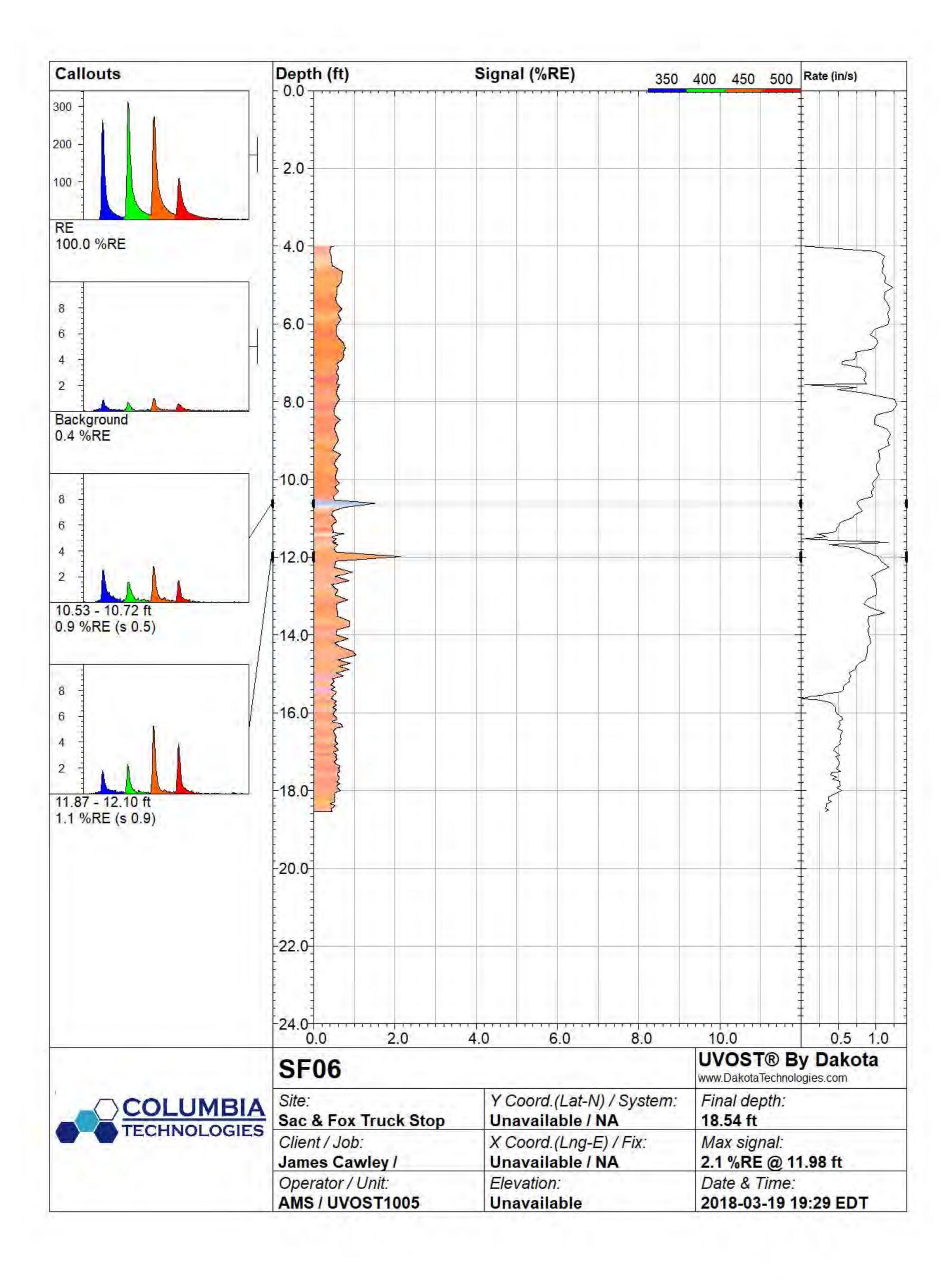
APPENDIX F – LIF/UVOST® Logs Collective Scale

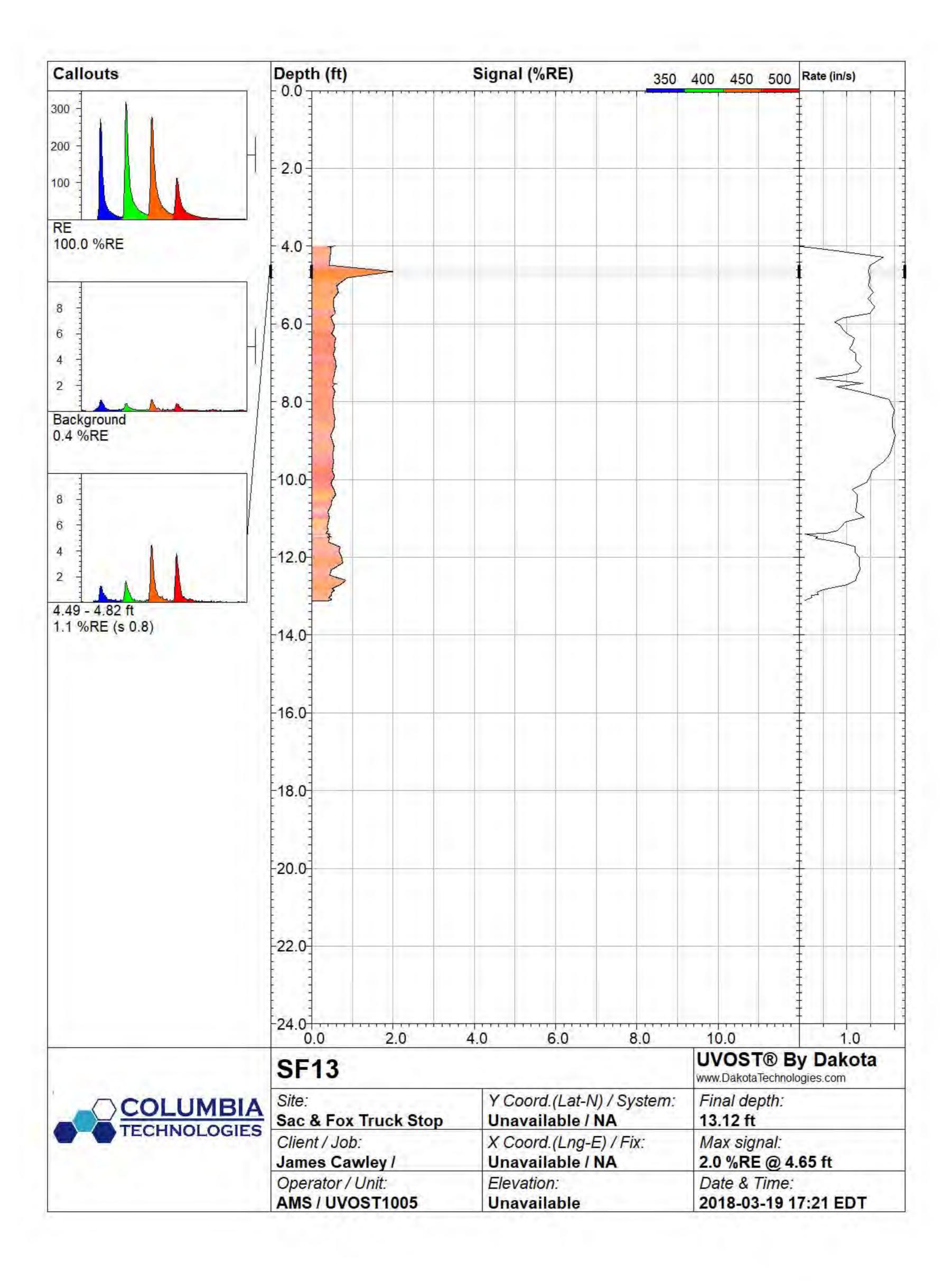


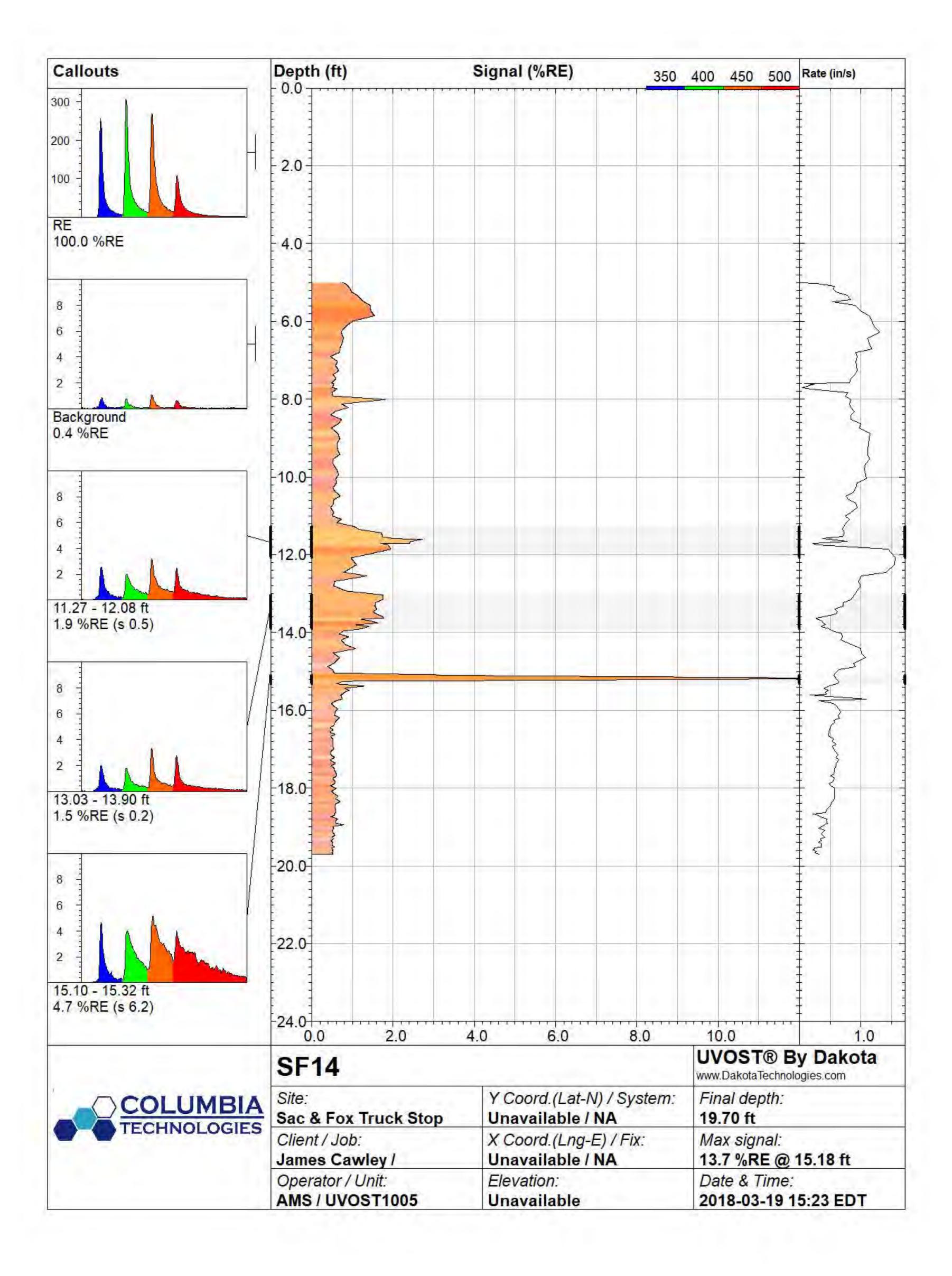


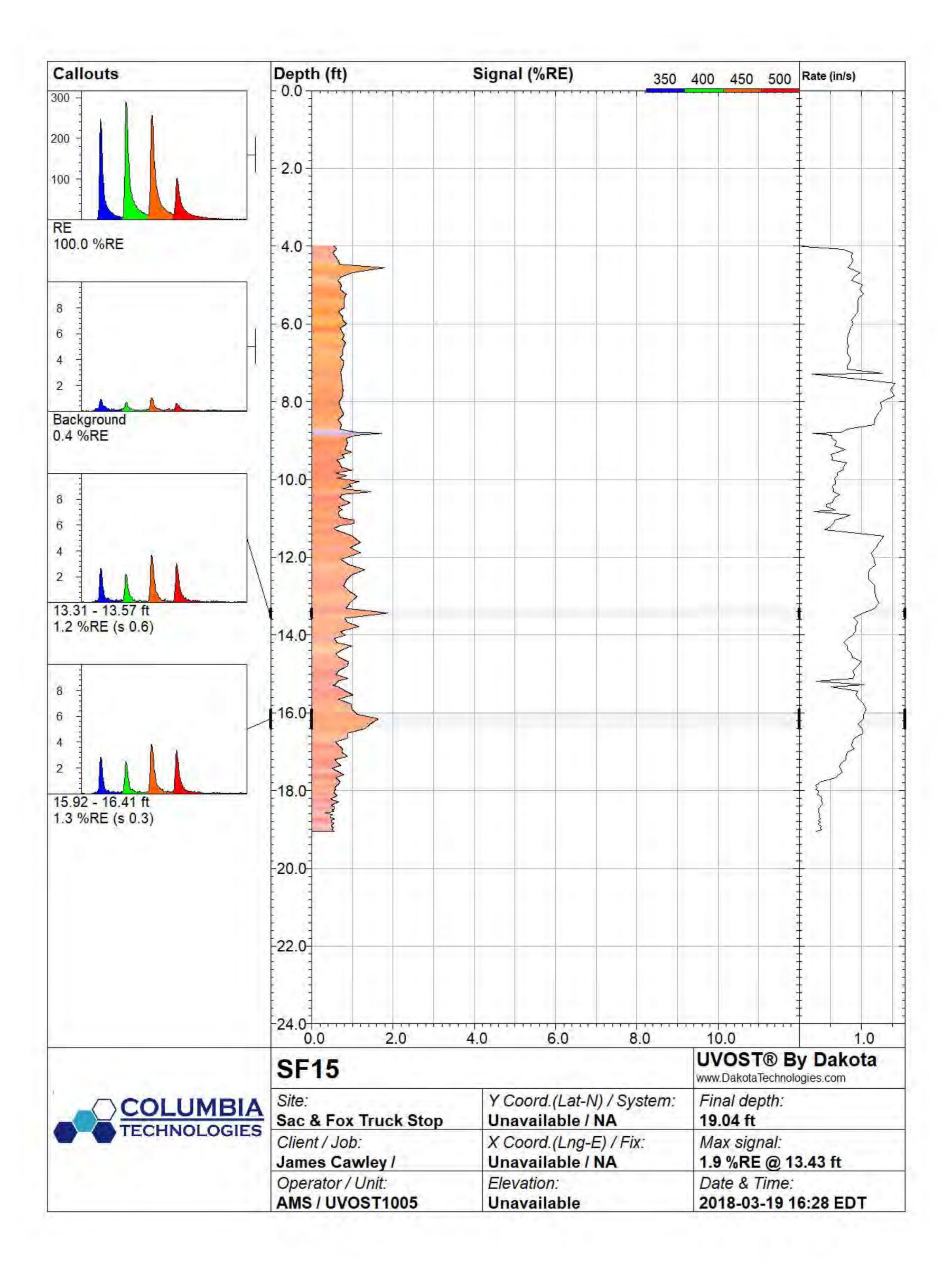




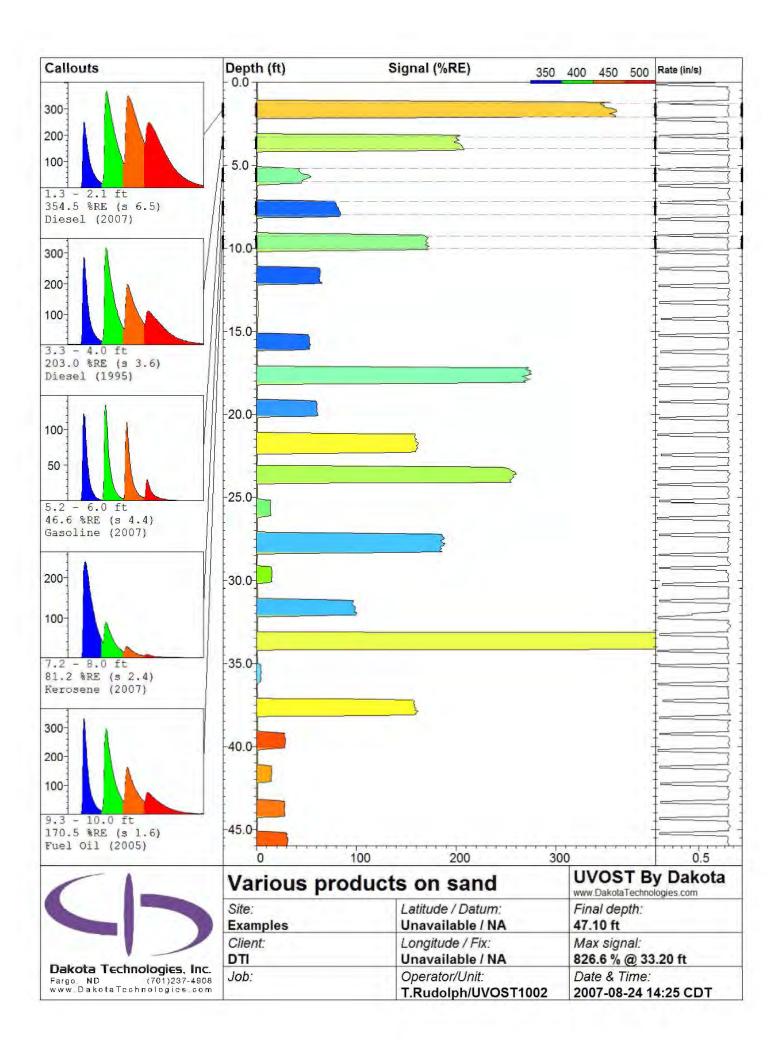


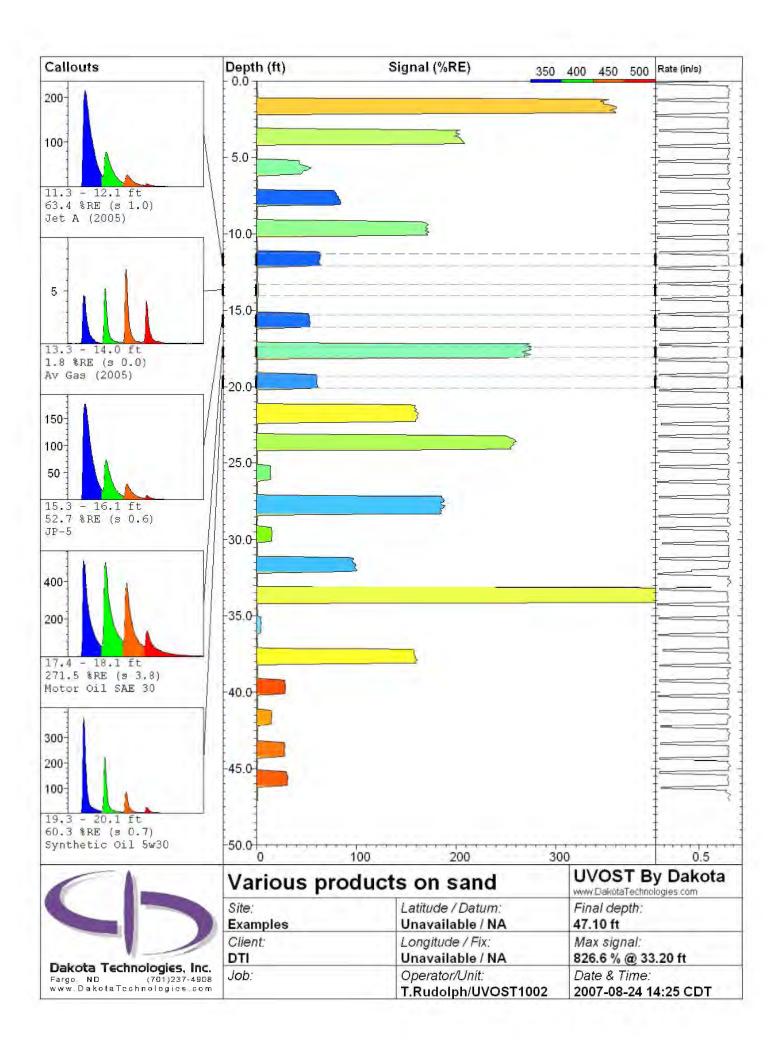


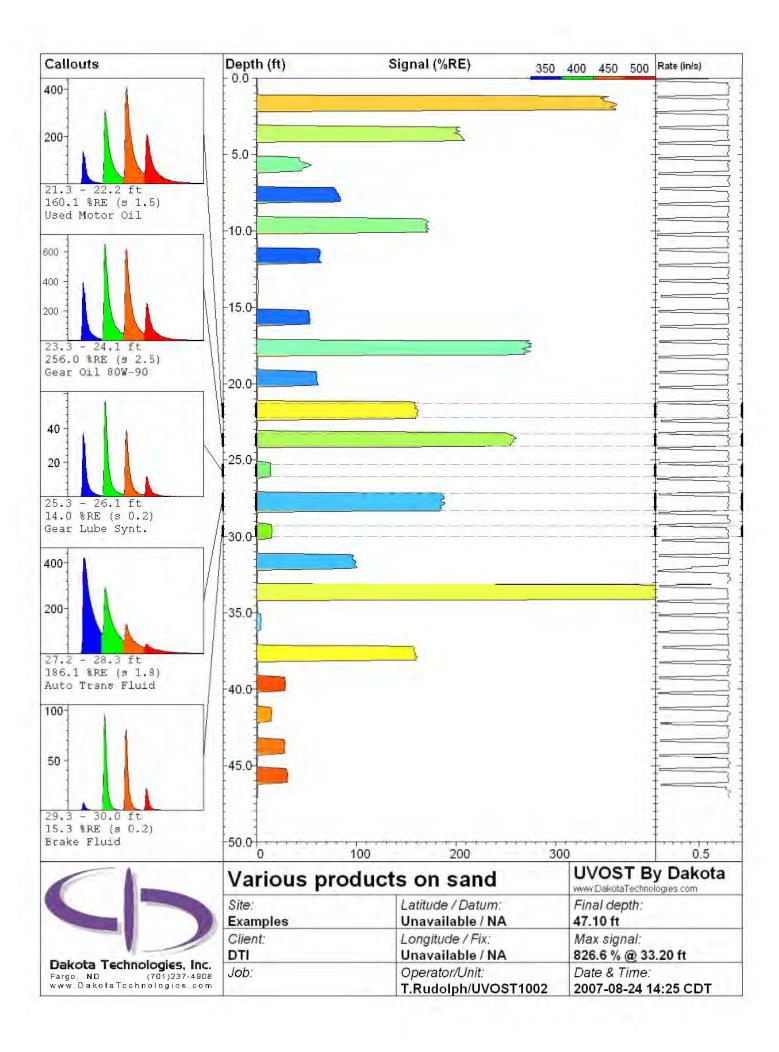


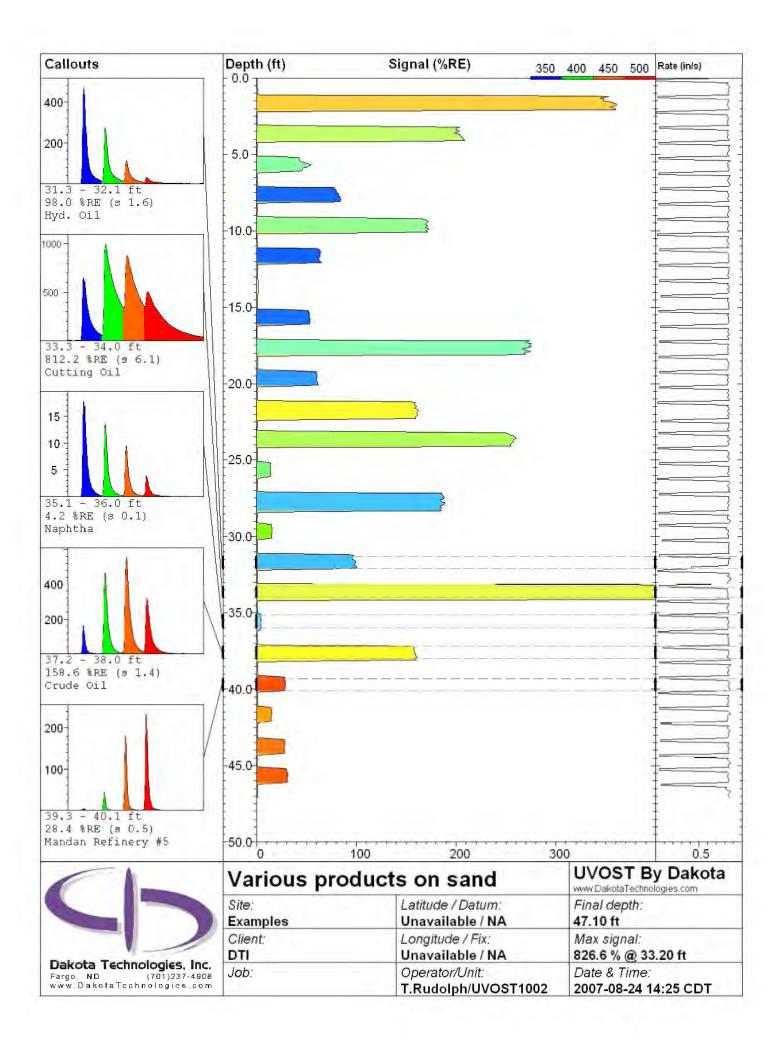


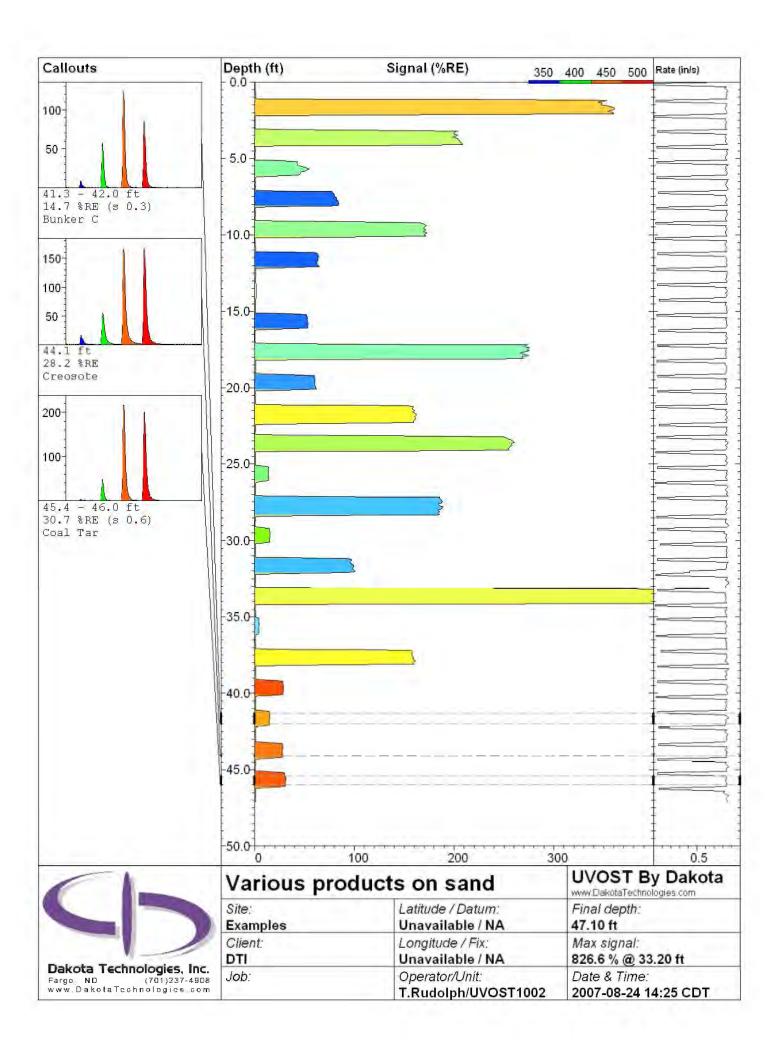
APPENDIX G – UVOST[®] Response to Various Saturated Products on Wet Sand











APPENDIX H – Analytical Lab Results Provided By ALS Environmental Laboratory



10450 Stancliff Rd. Suite 210 Houston, TX 77099 T: +1 281 530 5656

F: +1 281 530 5887

March 28, 2018

Scott Pieper Columbia Technologies LLC 1795 Cogswell Street Suite 101 Rockledge,, FL 32955

Work Order: **HS18031138**

Laboratory Results for: Sac + Fox

Dear Scott,

ALS Environmental received 26 sample(s) on Mar 23, 2018 for the analysis presented in the following report.

The analytical data provided relates directly to the samples received by ALS Environmental and for only the analyses requested. Results are expressed as "as received" unless otherwise noted.

QC sample results for this data met EPA or laboratory specifications except as noted in the Case Narrative or as noted with qualifiers in the QC batch information. Should this laboratory report need to be reproduced, it should be reproduced in full unless written approval has been obtained by ALS Environmental. Samples will be disposed in 30 days unless storage arrangements are made.

If you have any questions regarding this report, please feel free to call me.

Sincerely,

Generated By: JUMOKE.LAWAL

Corey Grandits
Project Manager

Project: Sac + Fox
Work Order: HS18031138

SAMPLE SUMMARY

Lab Samp ID	Client Sample ID	Matrix	TagNo	Collection Date	Date Received	Hold
HS18031138-01	SB01-11	Soil		22-Mar-2018 12:21	23-Mar-2018 08:50	
HS18031138-02	SB01-12	Soil		22-Mar-2018 12:26	23-Mar-2018 08:50	
HS18031138-03	SB01-13	Soil		22-Mar-2018 12:31	23-Mar-2018 08:50	
HS18031138-04	SB01-14	Soil		22-Mar-2018 12:36	23-Mar-2018 08:50	
HS18031138-05	SB01-15	Soil		22-Mar-2018 12:40	23-Mar-2018 08:50	
HS18031138-06	SB02-12	Soil		22-Mar-2018 12:55	23-Mar-2018 08:50	
HS18031138-07	SB02-14	Soil		22-Mar-2018 13:06	23-Mar-2018 08:50	
HS18031138-08	SB02-16	Soil		22-Mar-2018 15:20	23-Mar-2018 08:50	
HS18031138-09	SB03-20	Soil		22-Mar-2018 13:26	23-Mar-2018 08:50	
HS18031138-10	SB03-21	Soil		22-Mar-2018 13:33	23-Mar-2018 08:50	
HS18031138-11	SB03-22	Soil		22-Mar-2018 13:39	23-Mar-2018 08:50	
HS18031138-12	SB04-11	Soil		22-Mar-2018 14:05	23-Mar-2018 08:50	
HS18031138-13	SB04-12	Soil		22-Mar-2018 14:25	23-Mar-2018 08:50	
HS18031138-14	SB04-13	Soil		22-Mar-2018 14:10	23-Mar-2018 08:50	
HS18031138-15	SB04-14	Soil		22-Mar-2018 14:15	23-Mar-2018 08:50	
HS18031138-16	SB04-15	Soil		22-Mar-2018 14:20	23-Mar-2018 08:50	
HS18031138-17	SB04-16	Soil		22-Mar-2018 14:30	23-Mar-2018 08:50	
HS18031138-18	SB05-8	Soil		22-Mar-2018 15:10	23-Mar-2018 08:50	
HS18031138-19	SB05-9	Soil		22-Mar-2018 15:15	23-Mar-2018 08:50	
HS18031138-20	SB05-10	Soil		22-Mar-2018 14:47	23-Mar-2018 08:50	
HS18031138-21	SB05-11	Soil		22-Mar-2018 14:51	23-Mar-2018 08:50	
HS18031138-22	SB05-12	Soil		22-Mar-2018 14:55	23-Mar-2018 08:50	
HS18031138-23	SB05-13	Soil		22-Mar-2018 15:00	23-Mar-2018 08:50	
HS18031138-24	SB05-14	Soil		22-Mar-2018 15:06	23-Mar-2018 08:50	
HS18031138-25	SB05-15	Soil		22-Mar-2018 14:42	23-Mar-2018 08:50	
HS18031138-26	SB05-16	Soil		22-Mar-2018 14:45	23-Mar-2018 08:50	

ALS Group Houston, Corp

Client: Columbia Technologies LLC CASE NARRATIVE

Date: 28-Mar-18

Project: Sac + Fox
Work Order: HS18031138

GC Semivolatiles by Method TX1005

Batch ID: 126579,126588

• The test results meet requirements of the current NELAP standards, state requirements or programs where applicable.

Project: Sac + Fox Sample ID: SB01-11

Collection Date: 22-Mar-2018 12:21

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-01

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PF	R / 26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		38	mg/Kg	1	27-Mar-2018 00:31
>nC12 to nC28	ND		38	mg/Kg	1	27-Mar-2018 00:31
>nC28 to nC35	ND		38	mg/Kg	1	27-Mar-2018 00:31
Total Petroleum Hydrocarbon	ND		38	mg/Kg	1	27-Mar-2018 00:31
Surr: 2-Fluorobiphenyl	72.5		70-130	%REC	1	27-Mar-2018 00:31
Surr: Trifluoromethyl benzene	85.4		70-130	%REC	1	27-Mar-2018 00:31

Project: Sac + Fox Sample ID: SB01-12

Collection Date: 22-Mar-2018 12:26

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-02

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		40	mg/Kg	1	27-Mar-2018 01:00
>nC12 to nC28	ND		40	mg/Kg	1	27-Mar-2018 01:00
>nC28 to nC35	ND		40	mg/Kg	1	27-Mar-2018 01:00
Total Petroleum Hydrocarbon	ND		40	mg/Kg	1	27-Mar-2018 01:00
Surr: 2-Fluorobiphenyl	71.7		70-130	%REC	1	27-Mar-2018 01:00
Surr: Trifluoromethyl benzene	78.8		70-130	%REC	1	27-Mar-2018 01:00

Client: Columbia Technologies LLC

Project: Sac + Fox Sample ID: SB01-13

Collection Date: 22-Mar-2018 12:31

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-03

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		37	mg/Kg	1	27-Mar-2018 01:29
>nC12 to nC28	60		37	mg/Kg	1	27-Mar-2018 01:29
>nC28 to nC35	ND		37	mg/Kg	1	27-Mar-2018 01:29
Total Petroleum Hydrocarbon	60.0		37	mg/Kg	1	27-Mar-2018 01:29
Surr: 2-Fluorobiphenyl	70.5		70-130	%REC	1	27-Mar-2018 01:29
Surr: Trifluoromethyl benzene	78.7		70-130	%REC	1	27-Mar-2018 01:29

Project: Sac + Fox Sample ID: SB01-14

Collection Date: 22-Mar-2018 12:36

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-04

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005	LIMIT	Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	60		38	mg/Kg		27-Mar-2018 01:58
>nC12 to nC28	360		38	mg/Kg	1	27-Mar-2018 01:58
>nC28 to nC35	ND		38	mg/Kg	1	27-Mar-2018 01:58
Total Petroleum Hydrocarbon	420		38	mg/Kg	1	27-Mar-2018 01:58
Surr: 2-Fluorobiphenyl	87.4		70-130	%REC	1	27-Mar-2018 01:58
Surr: Trifluoromethyl benzene	90.7		70-130	%REC	1	27-Mar-2018 01:58

Project: Sac + Fox Sample ID: SB01-15

Collection Date: 22-Mar-2018 12:40

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-05

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	45		38	mg/Kg	1	27-Mar-2018 02:27
>nC12 to nC28	330		38	mg/Kg	1	27-Mar-2018 02:27
>nC28 to nC35	ND		38	mg/Kg	1	27-Mar-2018 02:27
Total Petroleum Hydrocarbon	375		38	mg/Kg	1	27-Mar-2018 02:27
Surr: 2-Fluorobiphenyl	83.3		70-130	%REC	1	27-Mar-2018 02:27
Surr: Trifluoromethyl benzene	83.0		70-130	%REC	1	27-Mar-2018 02:27

Project: Sac + Fox Sample ID: SB02-12

Collection Date: 22-Mar-2018 12:55

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-06

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	26-Mar-2018	Analyst: MBG
nC6 to nC12	190		39	mg/Kg	1	27-Mar-2018 02:56
>nC12 to nC28	ND		39	mg/Kg	1	27-Mar-2018 02:56
>nC28 to nC35	ND		39	mg/Kg	1	27-Mar-2018 02:56
Total Petroleum Hydrocarbon	190		39	mg/Kg	1	27-Mar-2018 02:56
Surr: 2-Fluorobiphenyl	70.3		70-130	%REC	1	27-Mar-2018 02:56
Surr: Trifluoromethyl benzene	92.2		70-130	%REC	1	27-Mar-2018 02:56

Project: Sac + Fox Sample ID: SB02-14

Collection Date: 22-Mar-2018 13:06

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-07

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	140		38	mg/Kg	1	26-Mar-2018 17:45
>nC12 to nC28	ND		38	mg/Kg	1	26-Mar-2018 17:45
>nC28 to nC35	ND		38	mg/Kg	1	26-Mar-2018 17:45
Total Petroleum Hydrocarbon	140		38	mg/Kg	1	26-Mar-2018 17:45
Surr: 2-Fluorobiphenyl	70.6		70-130	%REC	1	26-Mar-2018 17:45
Surr: Trifluoromethyl benzene	97.3		70-130	%REC	1	26-Mar-2018 17:45

Project: Sac + Fox Sample ID: SB02-16

Collection Date: 22-Mar-2018 15:20

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-08

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		39	mg/Kg	1	27-Mar-2018 03:25
>nC12 to nC28	ND		39	mg/Kg	1	27-Mar-2018 03:25
>nC28 to nC35	ND		39	mg/Kg	1	27-Mar-2018 03:25
Total Petroleum Hydrocarbon	ND		39	mg/Kg	1	27-Mar-2018 03:25
Surr: 2-Fluorobiphenyl	71.4		70-130	%REC	1	27-Mar-2018 03:25
Surr: Trifluoromethyl benzene	79.2		70-130	%REC	1	27-Mar-2018 03:25

Project: Sac + Fox Sample ID: SB03-20

Collection Date: 22-Mar-2018 13:26

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-09

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		38	mg/Kg	1	27-Mar-2018 03:55
>nC12 to nC28	ND		38	mg/Kg	1	27-Mar-2018 03:55
>nC28 to nC35	ND		38	mg/Kg	1	27-Mar-2018 03:55
Total Petroleum Hydrocarbon	ND		38	mg/Kg	1	27-Mar-2018 03:55
Surr: 2-Fluorobiphenyl	72.9		70-130	%REC	1	27-Mar-2018 03:55
Surr: Trifluoromethyl benzene	80.7		70-130	%REC	1	27-Mar-2018 03:55

Client: Columbia Technologies LLC

Project: Sac + Fox Sample ID: SB03-21

Collection Date: 22-Mar-2018 13:33

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-10

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		38	mg/Kg	1	27-Mar-2018 04:24
>nC12 to nC28	ND		38	mg/Kg	1	27-Mar-2018 04:24
>nC28 to nC35	ND		38	mg/Kg	1	27-Mar-2018 04:24
Total Petroleum Hydrocarbon	ND		38	mg/Kg	1	27-Mar-2018 04:24
Surr: 2-Fluorobiphenyl	72.1		70-130	%REC	1	27-Mar-2018 04:24
Surr: Trifluoromethyl benzene	71.8		70-130	%REC	1	27-Mar-2018 04:24

Project: Sac + Fox Sample ID: SB03-22

Collection Date: 22-Mar-2018 13:39

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-11

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PF	R / 26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		37	mg/Kg	1	27-Mar-2018 04:53
>nC12 to nC28	ND		37	mg/Kg	1	27-Mar-2018 04:53
>nC28 to nC35	ND		37	mg/Kg	1	27-Mar-2018 04:53
Total Petroleum Hydrocarbon	ND		37	mg/Kg	1	27-Mar-2018 04:53
Surr: 2-Fluorobiphenyl	78.9		70-130	%REC	1	27-Mar-2018 04:53
Surr: Trifluoromethyl benzene	90.3		70-130	%REC	1	27-Mar-2018 04:53

Client: Columbia Technologies LLC

Project: Sac + Fox Sample ID: SB04-11

Collection Date: 22-Mar-2018 14:05

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-12

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR /	26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		39	mg/Kg	1	27-Mar-2018 15:45
>nC12 to nC28	ND		39	mg/Kg	1	27-Mar-2018 15:45
>nC28 to nC35	ND		39	mg/Kg	1	27-Mar-2018 15:45
Total Petroleum Hydrocarbon	ND		39	mg/Kg	1	27-Mar-2018 15:45
Surr: 2-Fluorobiphenyl	75.4		70-130	%REC	1	27-Mar-2018 15:45
Surr: Trifluoromethyl benzene	85.1		70-130	%REC	1	27-Mar-2018 15:45

Project: Sac + Fox Sample ID: SB04-12

Collection Date: 22-Mar-2018 14:25

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-13

Matrix:Soil

			REPORT		DILUTION	DATE
ANALYSES	RESULT	QUAL	LIMIT	UNITS	FACTOR	ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		38	mg/Kg	1	27-Mar-2018 16:13
>nC12 to nC28	ND		38	mg/Kg	1	27-Mar-2018 16:13
>nC28 to nC35	ND		38	mg/Kg	1	27-Mar-2018 16:13
Total Petroleum Hydrocarbon	ND		38	mg/Kg	1	27-Mar-2018 16:13
Surr: 2-Fluorobiphenyl	73.3		70-130	%REC	1	27-Mar-2018 16:13
Surr: Trifluoromethyl benzene	84.4		70-130	%REC	1	27-Mar-2018 16:13

Project: Sac + Fox Sample ID: SB04-13

Collection Date: 22-Mar-2018 14:10

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-14

Matrix:Soil

					DILUTION	DATE
ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	FACTOR	ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		39	mg/Kg	1	27-Mar-2018 16:42
>nC12 to nC28	ND		39	mg/Kg	1	27-Mar-2018 16:42
>nC28 to nC35	ND		39	mg/Kg	1	27-Mar-2018 16:42
Total Petroleum Hydrocarbon	ND		39	mg/Kg	1	27-Mar-2018 16:42
Surr: 2-Fluorobiphenyl	73.0		70-130	%REC	1	27-Mar-2018 16:42
Surr: Trifluoromethyl benzene	82.4		70-130	%REC	1	27-Mar-2018 16:42

Project: Sac + Fox Sample ID: SB04-14

Collection Date: 22-Mar-2018 14:15

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-15

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		36	mg/Kg	1	27-Mar-2018 17:11
>nC12 to nC28	ND		36	mg/Kg	1	27-Mar-2018 17:11
>nC28 to nC35	ND		36	mg/Kg	1	27-Mar-2018 17:11
Total Petroleum Hydrocarbon	ND		36	mg/Kg	1	27-Mar-2018 17:11
Surr: 2-Fluorobiphenyl	77.4		70-130	%REC	1	27-Mar-2018 17:11
Surr: Trifluoromethyl benzene	88.1		70-130	%REC	1	27-Mar-2018 17:11

Project: Sac + Fox Sample ID: SB04-15

Collection Date: 22-Mar-2018 14:20

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-16

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		39	mg/Kg	1	27-Mar-2018 17:40
>nC12 to nC28	ND		39	mg/Kg	1	27-Mar-2018 17:40
>nC28 to nC35	ND		39	mg/Kg	1	27-Mar-2018 17:40
Total Petroleum Hydrocarbon	ND		39	mg/Kg	1	27-Mar-2018 17:40
Surr: 2-Fluorobiphenyl	75.2		70-130	%REC	1	27-Mar-2018 17:40
Surr: Trifluoromethyl benzene	86.2		70-130	%REC	1	27-Mar-2018 17:40

Project: Sac + Fox Sample ID: SB04-16

Collection Date: 22-Mar-2018 14:30

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-17

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		40	mg/Kg	1	27-Mar-2018 18:09
>nC12 to nC28	ND		40	mg/Kg	1	27-Mar-2018 18:09
>nC28 to nC35	ND		40	mg/Kg	1	27-Mar-2018 18:09
Total Petroleum Hydrocarbon	ND		40	mg/Kg	1	27-Mar-2018 18:09
Surr: 2-Fluorobiphenyl	76.1		70-130	%REC	1	27-Mar-2018 18:09
Surr: Trifluoromethyl benzene	85.6		70-130	%REC	1	27-Mar-2018 18:09

Project: Sac + Fox Sample ID: SB05-8

Collection Date: 22-Mar-2018 15:10

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-18

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MB0
nC6 to nC12	130		40	mg/Kg	1	27-Mar-2018 18:3
>nC12 to nC28	ND		40	mg/Kg	1	27-Mar-2018 18:3
>nC28 to nC35	ND		40	mg/Kg	1	27-Mar-2018 18:3
Total Petroleum Hydrocarbon	130		40	mg/Kg	1	27-Mar-2018 18:3
Surr: 2-Fluorobiphenyl	76.6		70-130	%REC	1	27-Mar-2018 18:3
Surr: Trifluoromethyl benzene	86.1		70-130	%REC	1	27-Mar-2018 18:3

Project: Sac + Fox Sample ID: SB05-9

Collection Date: 22-Mar-2018 15:15

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-19

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	110		38	mg/Kg	1	27-Mar-2018 19:07
>nC12 to nC28	ND		38	mg/Kg	1	27-Mar-2018 19:07
>nC28 to nC35	ND		38	mg/Kg	1	27-Mar-2018 19:07
Total Petroleum Hydrocarbon	110		38	mg/Kg	1	27-Mar-2018 19:07
Surr: 2-Fluorobiphenyl	80.0		70-130	%REC	1	27-Mar-2018 19:07
Surr: Trifluoromethyl benzene	87.2		70-130	%REC	1	27-Mar-2018 19:07

Project: Sac + Fox Sample ID: SB05-10

Collection Date: 22-Mar-2018 14:47

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-20

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		38	mg/Kg	1	27-Mar-2018 19:36
>nC12 to nC28	ND		38	mg/Kg	1	27-Mar-2018 19:36
>nC28 to nC35	ND		38	mg/Kg	1	27-Mar-2018 19:36
Total Petroleum Hydrocarbon	ND		38	mg/Kg	1	27-Mar-2018 19:36
Surr: 2-Fluorobiphenyl	75.8		70-130	%REC	1	27-Mar-2018 19:36
Surr: Trifluoromethyl benzene	86.9		70-130	%REC	1	27-Mar-2018 19:36

Client: Columbia Technologies LLC

Project: Sac + Fox Sample ID: SB05-11

Collection Date: 22-Mar-2018 14:51

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-21

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		39	mg/Kg	1	27-Mar-2018 20:05
>nC12 to nC28	ND		39	mg/Kg	1	27-Mar-2018 20:05
>nC28 to nC35	ND		39	mg/Kg	1	27-Mar-2018 20:05
Total Petroleum Hydrocarbon	ND		39	mg/Kg	1	27-Mar-2018 20:05
Surr: 2-Fluorobiphenyl	78.8		70-130	%REC	1	27-Mar-2018 20:05
Surr: Trifluoromethyl benzene	89.5		70-130	%REC	1	27-Mar-2018 20:05

Client: Columbia Technologies LLC

Project: Sac + Fox Sample ID: SB05-12

Collection Date: 22-Mar-2018 14:55

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-22

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		38	mg/Kg	1	27-Mar-2018 10:55
>nC12 to nC28	ND		38	mg/Kg	1	27-Mar-2018 10:55
>nC28 to nC35	ND		38	mg/Kg	1	27-Mar-2018 10:55
Total Petroleum Hydrocarbon	ND		38	mg/Kg	1	27-Mar-2018 10:55
Surr: 2-Fluorobiphenyl	74.0		70-130	%REC	1	27-Mar-2018 10:55
Surr: Trifluoromethyl benzene	84.2		70-130	%REC	1	27-Mar-2018 10:55

Client: Columbia Technologies LLC

Project: Sac + Fox Sample ID: SB05-13

Collection Date: 22-Mar-2018 15:00

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-23

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		38	mg/Kg	1	27-Mar-2018 20:34
>nC12 to nC28	ND		38	mg/Kg	1	27-Mar-2018 20:34
>nC28 to nC35	ND		38	mg/Kg	1	27-Mar-2018 20:34
Total Petroleum Hydrocarbon	ND		38	mg/Kg	1	27-Mar-2018 20:34
Surr: 2-Fluorobiphenyl	74.6		70-130	%REC	1	27-Mar-2018 20:34
Surr: Trifluoromethyl benzene	83.8		70-130	%REC	1	27-Mar-2018 20:34

Project: Sac + Fox Sample ID: SB05-14

Collection Date: 22-Mar-2018 15:06

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-24

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		39	mg/Kg	1	27-Mar-2018 21:02
>nC12 to nC28	ND		39	mg/Kg	1	27-Mar-2018 21:02
>nC28 to nC35	ND		39	mg/Kg	1	27-Mar-2018 21:02
Total Petroleum Hydrocarbon	ND		39	mg/Kg	1	27-Mar-2018 21:02
Surr: 2-Fluorobiphenyl	75.4		70-130	%REC	1	27-Mar-2018 21:02
Surr: Trifluoromethyl benzene	83.8		70-130	%REC	1	27-Mar-2018 21:02

Client: Columbia Technologies LLC

Project: Sac + Fox Sample ID: SB05-15

Collection Date: 22-Mar-2018 14:42

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-25

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		39	mg/Kg	1	27-Mar-2018 21:31
>nC12 to nC28	ND		39	mg/Kg	1	27-Mar-2018 21:31
>nC28 to nC35	ND		39	mg/Kg	1	27-Mar-2018 21:31
Total Petroleum Hydrocarbon	ND		39	mg/Kg	1	27-Mar-2018 21:31
Surr: 2-Fluorobiphenyl	72.6		70-130	%REC	1	27-Mar-2018 21:31
Surr: Trifluoromethyl benzene	84.3		70-130	%REC	1	27-Mar-2018 21:31

Project: Sac + Fox Sample ID: SB05-16

Collection Date: 22-Mar-2018 14:45

ANALYTICAL REPORT

WorkOrder:HS18031138 Lab ID:HS18031138-26

Matrix:Soil

ANALYSES	RESULT	QUAL	REPORT LIMIT	UNITS	DILUTION FACTOR	DATE ANALYZED
TEXAS TPH BY TX1005		Method:TX1005		Prep:TX1005PR	/ 26-Mar-2018	Analyst: MBG
nC6 to nC12	ND		38	mg/Kg	1	27-Mar-2018 22:59
>nC12 to nC28	ND		38	mg/Kg	1	27-Mar-2018 22:59
>nC28 to nC35	ND		38	mg/Kg	1	27-Mar-2018 22:59
Total Petroleum Hydrocarbon	ND		38	mg/Kg	1	27-Mar-2018 22:59
Surr: 2-Fluorobiphenyl	74.0		70-130	%REC	1	27-Mar-2018 22:59
Surr: Trifluoromethyl benzene	84.6		70-130	%REC	1	27-Mar-2018 22:59

WEIGHT LOG

Client: Columbia Technologies LLC

Project: Sac + Fox **WorkOrder:** HS18031138

Batch ID : 126579	Metho	d: TEXAS	TPH BY TX10	005	Prep: TX 1005_S PR
SampID	Container	Sample Wt/Vol	Final Volume	Prep Factor	
HS18031138-01	1	13.28	10 (mL)	0.753	
HS18031138-02	1	12.42	10 (mL)	0.8052	
HS18031138-03	1	13.38	10 (mL)	0.7474	
HS18031138-04	1	13.07	10 (mL)	0.7651	
HS18031138-05	1	13.17	10 (mL)	0.7593	
HS18031138-06	1	12.95	10 (mL)	0.7722	
HS18031138-07	1	13.24	10 (mL)	0.7553	
HS18031138-08	1	12.73	10 (mL)	0.7855	
HS18031138-09	1	13.06	10 (mL)	0.7657	
HS18031138-10	1	13.33	10 (mL)	0.7502	
HS18031138-11	1	13.68	10 (mL)	0.731	

Batch ID : 126588	Metho	d: TEXAS	TPH BY TX10	005	Prep: TX 1005_S PR
SampID	Container	Sample Wt/Vol	Final Volume	Prep Factor	
HS18031138-12	1	12.81	10 (mL)	0.7806	
HS18031138-13	1	13.13	10 (mL)	0.7616	
HS18031138-14	1	12.79	10 (mL)	0.7819	
HS18031138-15	1	13.8	10 (mL)	0.7246	
HS18031138-16	1	12.85	10 (mL)	0.7782	
HS18031138-17	1	12.62	10 (mL)	0.7924	
HS18031138-18	1	12.49	10 (mL)	0.8006	
HS18031138-19	1	13.14	10 (mL)	0.761	
HS18031138-20	1	13.13	10 (mL)	0.7616	
HS18031138-21	1	12.92	10 (mL)	0.774	
HS18031138-22	1	13.3	10 (mL)	0.7519	
HS18031138-23	1	13.28	10 (mL)	0.753	
HS18031138-24	1	12.78	10 (mL)	0.7825	
HS18031138-25	1	12.69	10 (mL)	0.788	
HS18031138-26	1	13.32	10 (mL)	0.7508	

Project: Sac + Fox DATES REPORT

WorkOrder: HS18031138

Sample ID	Client Sar	mp ID Collection Date	TCLP Date	Prep Date	Analysis Date	DF
Batch ID 12657	9	Test Name: TEXAS TPH BY TX1005		Matrix: S	Soil	
HS18031138-01	SB01-11	22 Mar 2018 12:21		26 Mar 2018 09:48	27 Mar 2018 00:31	1
HS18031138-02	SB01-12	22 Mar 2018 12:26		26 Mar 2018 09:48	27 Mar 2018 01:00	1
HS18031138-03	SB01-13	22 Mar 2018 12:31		26 Mar 2018 09:48	27 Mar 2018 01:29	1
HS18031138-04	SB01-14	22 Mar 2018 12:36		26 Mar 2018 09:48	27 Mar 2018 01:58	1
HS18031138-05	SB01-15	22 Mar 2018 12:40		26 Mar 2018 09:48	27 Mar 2018 02:27	1
HS18031138-06	SB02-12	22 Mar 2018 12:55		26 Mar 2018 09:48	27 Mar 2018 02:56	1
HS18031138-07	SB02-14	22 Mar 2018 13:06		26 Mar 2018 09:48	26 Mar 2018 17:45	1
HS18031138-08	SB02-16	22 Mar 2018 15:20		26 Mar 2018 09:48	27 Mar 2018 03:25	1
HS18031138-09	SB03-20	22 Mar 2018 13:26		26 Mar 2018 09:48	27 Mar 2018 03:55	1
HS18031138-10	SB03-21	22 Mar 2018 13:33		26 Mar 2018 09:48	27 Mar 2018 04:24	1
HS18031138-11	SB03-22	22 Mar 2018 13:39		26 Mar 2018 09:48	27 Mar 2018 04:53	1
Batch ID 12658	8	Test Name: TEXAS TPH BY TX1005		Matrix: S	Soil	
HS18031138-12	SB04-11	22 Mar 2018 14:05		26 Mar 2018 11:27	27 Mar 2018 15:45	1
HS18031138-13	SB04-12	22 Mar 2018 14:25		26 Mar 2018 11:27	27 Mar 2018 16:13	1
HS18031138-14	SB04-13	22 Mar 2018 14:10		26 Mar 2018 11:27	27 Mar 2018 16:42	1
HS18031138-15	SB04-14	22 Mar 2018 14:15		26 Mar 2018 11:27	27 Mar 2018 17:11	1
HS18031138-16	SB04-15	22 Mar 2018 14:20		26 Mar 2018 11:27	27 Mar 2018 17:40	1
HS18031138-17	SB04-16	22 Mar 2018 14:30		26 Mar 2018 11:27	27 Mar 2018 18:09	1
HS18031138-18	SB05-8	22 Mar 2018 15:10		26 Mar 2018 11:27	27 Mar 2018 18:38	1
HS18031138-19	SB05-9	22 Mar 2018 15:15		26 Mar 2018 11:27	27 Mar 2018 19:07	1
HS18031138-20	SB05-10	22 Mar 2018 14:47		26 Mar 2018 11:27	27 Mar 2018 19:36	1
HS18031138-21	SB05-11	22 Mar 2018 14:51		26 Mar 2018 11:27	27 Mar 2018 20:05	1
HS18031138-22	SB05-12	22 Mar 2018 14:55		26 Mar 2018 11:27	27 Mar 2018 10:55	1
HS18031138-23	SB05-13	22 Mar 2018 15:00		26 Mar 2018 11:27	27 Mar 2018 20:34	1
HS18031138-24	SB05-14	22 Mar 2018 15:06		26 Mar 2018 11:27	27 Mar 2018 21:02	1
HS18031138-25	SB05-15	22 Mar 2018 14:42		26 Mar 2018 11:27	27 Mar 2018 21:31	1
HS18031138-26	SB05-16	22 Mar 2018 14:45		26 Mar 2018 11:27	27 Mar 2018 22:59	1

Client: Columbia Technologies LLC

Project: Sac + Fox WorkOrder: HS18031138

QC BATCH REPORT

nC6 to nC12	Batch ID: 126579			Instrument:	FID-13		Metho	od: TX1005	i	
Analyte Result PQL SPK Val SPK Ref Value REC Control Value RPD Ref RPD Ref Value Ref Value Ref Value RPD Ref RPD		Sample ID:	MBLK-126579	Pun ID: FID-1				•		
ND 50 SeqNo: 4489720 SeqNo: 4489			.			SPK Ref		Control	RPD Ref	RPD
>nC12 to nC28	Analyte		Result	PQL	SPK Val	Value	%REC	Limit	Value	%RPD Limit Qua
ND 50 Sur: 2-Fluorobiphenyl 17.72 0 25 0 70.9 70 - 130	nC6 to nC12		ND	50						
Total Petroleum Hydrocarbon ND 50 Surr. 2-Fluorobiphenyl 17.72 0 25 0 70.9 70 - 130 Surr. 2-Fluorobiphenyl 19.04 0 25 0 76.2 70 - 130 Surr. 2-Fluorobiphenyl 19.04 0 25 0 76.2 70 - 130 Surr. 2-Fluorobiphenyl 19.04 0 25 0 76.2 70 - 130 Section 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19.04 19	>nC12 to nC28		ND	50						
Surr: 2-Fluorobiphenyl 17.72 0 25 0 70.9 70 - 130	>nC28 to nC35		ND	50						
LCS	Total Petroleum Hyd	rocarbon	ND	50						
Client ID: Run ID: FID-13_313166 SeqNo: 4489720 PrepDate: 26-Mar-2018 16:47	Surr: 2-Fluorobiphen	ıyl	17.72	0	25	0	70.9	70 - 130		
Result PQL SPK Val SPK Val SPK Val SPK Ref	Surr: Trifluoromethyl	benzene	19.04	0	25	0	76.2	70 - 130		
Analyte Result PQL SPK Val Value Va	LCS	Sample ID:	LCS-126579		Units:	mg/Kg	Ana	alysis Date:	26-Mar-2018	16:47
Analyte Result PQL SPK Val Value %REC Limit Value %RPD Limit Concern	Client ID:			Run ID: FID-1	3_313166	SeqNo: 4	489720	PrepDate:	26-Mar-2018	DF: 1
NCC12 to nC28 244.4 50 250 0 97.8 75 - 125 Surr: 2-Fluorobiphenyl 19.85 0 25 0 79.4 70 - 130 LCSD Sample ID: LCSD-126579 Units: mg/Kg Analysis Date: 26-Mar-2018 17:16 Client ID: Run ID: FID-13_313166 SeqNo: 4489721 PrepDate: 26-Mar-2018 DF: 1 Analyte Result PQL SPK Val Value %REC Control Limit RPD Ref RPD Ref Value RPD Ref RPD Ref NepD Limit Control RPD Ref RPD Ref NepD Ref	Analyte		Result	PQL	SPK Val		%REC			RPD %RPD Limit Qua
Surr: 2-Fluorobiphenyl 19.85 0 25 0 79.4 70 - 130 Surr: Trifluoromethyl benzene 24.99 0 25 0 100.0 70 - 130 LCSD Sample ID: LCSD-126579 Units: mg/Kg Analysis Date: 26-Mar-2018 17:16 Client ID: Run ID: FID-13_313166 SeqNo: 4489721 PrepDate: 26-Mar-2018 DF: 1 Analyte Result PQL SPK Val SPK Ref AREC Control Limit RPD Ref Res 20 Res 20 8.88 20 20 Res 24.99 8.88 20 20 MS <td>nC6 to nC12</td> <td></td> <td>248.9</td> <td>50</td> <td>250</td> <td>0</td> <td>99.6</td> <td>75 - 125</td> <td></td> <td></td>	nC6 to nC12		248.9	50	250	0	99.6	75 - 125		
Control Cont	>nC12 to nC28		244.4	50	250	0	97.8	75 - 125		
LCSD Sample ID: LCSD-126579 Units: mg/Kg Analysis Date: 26-Mar-2018 17:16 Client ID: Run ID: FID-13_313166 SeqNo: 4489721 PrepDate: 26-Mar-2018 DF: 1 Analyte Result PQL SPK Val SPK Ref Value REC Control Limit RPD Ref RPD Ref RPD Ref RPD Ref RPD Limit Control Rep Result RPD Ref RPD Ref RPD Ref Value RPD Ref RPD Result Units: mg/Kg Analysis Date: 26-Mar-2018 18:14 Client ID: SB02-14 Run ID: FID-13_313166 SeqNo: 4489753 PrepDate: 26-Mar-2018 18:14 Client ID: SB02-14 Run ID: FID-13_313166 SeqNo: 4489753 PrepDate: 26-Mar-2018 18:14 Client ID: SB02-14 Result PQL SPK Val Value REC Control RPD Ref RP	Surr: 2-Fluorobiphen	ıyl	19.85	0	25	0	79.4	70 - 130		
Client ID: Run ID: FID-13_313166 SeqNo: 4489721 PrepDate: 26-Mar-2018 DF: 1 Analyte Result PQL SPK Val SPK Ref Value %REC Control RPD Ref Value %RPD Limit Concloration C2	Surr: Trifluoromethyl	benzene	24.99	0	25	0	100.0	70 - 130		
Analyte Result PQL SPK Val SPK Ref Value Control Limit RPD Ref Value RPD Limit Control Value RPD Limit Control Value RPD Ref Value RPD Limit Control Value RPD Ref Value RPD Limit Control Value RPD Ref Ref RPD Ref	LCSD	Sample ID:	LCSD-126579		Units:	mg/Kg	Ana	alysis Date:	26-Mar-2018	17:16
Analyte Result PQL SPK Val Value %REC Limit Value %RPD Limit Concept C	Client ID:			Run ID: FID-1	3_313166	SeqNo: 4	489721	PrepDate:	26-Mar-2018	DF: 1
NC12 to nC28 268.4 50 250 0 107 75 - 125 244.4 9.34 20 Surr: 2-Fluorobiphenyl 21.85 0 25 0 87.4 70 - 130 19.85 9.62 20 Surr: Trifluoromethyl benzene 27.31 0 25 0 109 70 - 130 24.99 8.88 20 MS Sample ID: HS18031138-07MS Units: mg/Kg Analysis Date: 26-Mar-2018 18:14 Client ID: SB02-14 Run ID: FID-13_313166 SeqNo: 4489753 PrepDate: 26-Mar-2018 DF: 1 Analyte Result PQL SPK Val Value %REC Control RPD Ref RPD nC6 to nC12 295.6 38 191.6 142.8 79.8 75 - 125 >nC12 to nC28 227.9 38 191.6 0 80.0 70 - 130 Surr: 2-Fluorobiphenyl 15.32 0 19.16 0 80.0 70 - 130	Analyte		Result	PQL	SPK Val		%REC			RPD %RPD Limit Qua
Surr: 2-Fluorobiphenyl 21.85 0 25 0 87.4 70 - 130 19.85 9.62 20 Surr: Trifluoromethyl benzene 27.31 0 25 0 109 70 - 130 24.99 8.88 20 MS Sample ID: HS18031138-07MS Units: mg/Kg Analysis Date: 26-Mar-2018 18:14 Client ID: SB02-14 Run ID: FID-13_313166 SeqNo: 4489753 PrepDate: 26-Mar-2018 DF: 1 SPK Ref Control RPD Ref RPD Analyte Result PQL SPK Val Value %REC Limit Value %RPD Limit Control PnC12 to nC28 227.9 38 191.6 142.8 79.8 75 - 125 Surr: 2-Fluorobiphenyl 15.32 0 19.16 0 80.0 70 - 130	nC6 to nC12		278.4	50	250	0	111	75 - 125	248.9	11.2 20
Surr: Trifluoromethyl benzene 27.31 0 25 0 109 70 - 130 24.99 8.88 20 MS Sample ID: HS18031138-07MS Units: mg/Kg Analysis Date: 26-Mar-2018 18:14 Client ID: SB02-14 Run ID: FID-13_313166 SeqNo: 4489753 PrepDate: 26-Mar-2018 DF: 1 SPK Ref Control RPD Ref RPD Ref RPD Analyte Result PQL SPK Val Value %REC Limit Value %RPD Limit Of nC6 to nC12 295.6 38 191.6 142.8 79.8 75 - 125 >nC12 to nC28 227.9 38 191.6 0 119 75 - 125 Surr: 2-Fluorobiphenyl 15.32 0 19.16 0 80.0 70 - 130	>nC12 to nC28		268.4	50	250	0	107	75 - 125	244.4	9.34 20
MS Sample ID: HS18031138-07MS Units: mg/Kg Analysis Date: 26-Mar-2018 18:14 Client ID: SB02-14 Run ID: FID-13_313166 SeqNo: 4489753 PrepDate: 26-Mar-2018 DF: 1 SPK Ref SPK Ref Control RPD Ref RPD Ref RPD Analyte Result PQL SPK Val Value %REC Limit Value %RPD Limit nC6 to nC12 295.6 38 191.6 142.8 79.8 75 - 125 >nC12 to nC28 227.9 38 191.6 0 119 75 - 125 Surr: 2-Fluorobiphenyl 15.32 0 19.16 0 80.0 70 - 130	Surr: 2-Fluorobiphen	ıyl	21.85	0	25	0	87.4	70 - 130	19.85	9.62 20
Client ID: SB02-14 Run ID: FID-13_313166 SeqNo: 4489753 PrepDate: 26-Mar-2018 DF: 1 Analyte Result PQL SPK Val Value %REC Control Limit RPD Ref RPD Limit Control Value NRPD Limit Control Value NRPD Limit Control Value NRPD Ref RPD Limit Control Value NRPD Limit Control Value NRPD Limit Control Value NRPD Ref RPD Ref RPD Limit Control Value NRPD Ref RPD Ref RPD Ref RPD Ref RPD Ref RPD Ref RPD Ref Ref RPD Ref	Surr: Trifluoromethyl	benzene	27.31	0	25	0	109	70 - 130	24.99	8.88 20
Analyte Result PQL SPK Val Value %REC Control RPD Ref RPD Limit Control NC6 to nC12 295.6 38 191.6 142.8 79.8 75 - 125 PnC12 to nC28 227.9 38 191.6 0 119 75 - 125 Surr: 2-Fluorobiphenyl 15.32 0 19.16 0 80.0 70 - 130	MS S	Sample ID:	HS18031138-07	MS	Units:	mg/Kg	Ana	alysis Date:	26-Mar-2018	18:14
Analyte Result PQL SPK Val Value %REC Limit Value %RPD Limit Of the Control of the Con	Client ID: SB02-14	4		Run ID: FID-1	3_313166	SeqNo: 4	489753	PrepDate:	26-Mar-2018	DF: 1
>nC12 to nC28 227.9 38 191.6 0 119 75 - 125 Surr: 2-Fluorobiphenyl 15.32 0 19.16 0 80.0 70 - 130	Analyte		Result	PQL	SPK Val		%REC			RPD %RPD Limit Qua
Surr: 2-Fluorobiphenyl 15.32 0 19.16 0 80.0 70 - 130	nC6 to nC12		295.6	38	191.6	142.8	79.8	75 - 125		
	>nC12 to nC28		227.9	38	191.6	0	119	75 - 125		
	Surr: 2-Fluorobiphen	nyl	15.32	0	19.16	0	80.0	70 - 130		
	Surr: Trifluoromethyl	benzene	20.2	0	19.16	0	105	70 - 130		

Client: Columbia Technologies LLC

Project: Sac + Fox WorkOrder: HS18031138

QC BATCH REPORT

Batch ID: 126579		ı	Instrument:	FID-13		Metho	d: TX1005			
MSD S	ample ID:	HS18031138-07M	SD	Units:	mg/Kg	Ana	alysis Date:	26-Mar-2018	18:43	77
Client ID: SB02-14		R	un ID: FID-13	_313166	SeqNo: 4	489724	PrepDate:	26-Mar-2018	DF: 1	
Analyte		Result	PQL	SPK Val	SPK Ref Value	%REC	Control Limit		R %RPD Li	PD mit Qua
nC6 to nC12		301.8	38	191.1	142.8	83.2	75 - 125	295.6	2.07	20
>nC12 to nC28		213.4	38	191.1	0	112	75 - 125	227.9	6.57	20
Surr: 2-Fluorobipheny	1	16.09	0	19.11	0	84.2	70 - 130	15.32	4.92	20
Surr: Trifluoromethyl l	benzene	20.45	0	19.11	0	107	70 - 130	20.2	1.23	20
The following samples v	vere analyze	HS18	031138-01 031138-05 031138-09	HS1803113 HS1803113 HS1803113	38-06	HS180311: HS180311: HS180311:	38-07	HS18031138-0		1

Client: Columbia Technologies LLC

Project: Sac + Fox WorkOrder: HS18031138

QC BATCH REPORT

Batch ID: 126588		Instrument:	FID-10		Metho	od: TX1005		
MBLK Sample	e ID: MBLK-126588		Units:	mg/Kg	Ana	alysis Date:	27-Mar-2018	09:28
Client ID:		Run ID: FID-	10_313250	SeqNo: 4	491753	PrepDate:	26-Mar-2018	DF: 1
Analyte	Result	PQL	SPK Val	SPK Ref Value	%REC	Control Limit	RPD Ref Value	RPD %RPD Limit Qua
nC6 to nC12	ND	50						
>nC12 to nC28	ND	50						
>nC28 to nC35	ND	50						
Total Petroleum Hydrocarb	on ND	50						
Surr: 2-Fluorobiphenyl	20.87	0	25	0	83.5	70 - 130		
Surr: Trifluoromethyl benze	ne 22.89	0	25	0	91.5	70 - 130		
LCS Sample	e ID: LCS-126588		Units:	mg/Kg	Ana	alysis Date:	27-Mar-2018	09:57
Client ID:		Run ID: FID-	10_313250	SeqNo: 4	491754	PrepDate:	26-Mar-2018	DF: 1
Analyte	Result	PQL	SPK Val	SPK Ref Value	%REC	Control Limit		RPD %RPD Limit Qua
nC6 to nC12	244.5	50	250	0	97.8	75 - 125		
>nC12 to nC28	250.1	50	250	0	100	75 - 125		
Surr: 2-Fluorobiphenyl	20.29	0	25	0	81.2	70 - 130		
Surr: Trifluoromethyl benze	ne 24.63	0	25	0	98.5	70 - 130		
LCSD Sample	e ID: LCSD-126588		Units:	mg/Kg	Ana	alysis Date:	27-Mar-2018	10:26
Client ID:		Run ID: FID-	10_313250	SeqNo: 4	491755	PrepDate:	26-Mar-2018	DF: 1
Analyte	Result	PQL	SPK Val	SPK Ref Value	%REC	Control Limit		RPD %RPD Limit Qua
nC6 to nC12	240.8	50	250	0	96.3	75 - 125	244.5	1.55 20
>nC12 to nC28	247.9	50	250	0	99.2	75 - 125	250.1	0.876 20
Surr: 2-Fluorobiphenyl	19.54	0	25	0	78.2	70 - 130	20.29	3.76 20
Surr: Trifluoromethyl benze	ne 23.9	0	25	0	95.6	70 - 130	24.63	3.01 20
MS Sample	e ID: HS18031138-22	MS	Units:	mg/Kg	Ana	alysis Date:	27-Mar-2018	11:24
Client ID: SB05-12		Run ID: FID-	10_313250	SeqNo: 4	491757	PrepDate:	26-Mar-2018	DF: 1
Analyte	Result	PQL	SPK Val	SPK Ref Value	%REC	Control Limit	RPD Ref Value	RPD %RPD Limit Qua
nC6 to nC12	223.9	39	194.7	0	115	75 - 125		
>nC12 to nC28	209.2		194.7	0	107	75 - 125		
Surr: 2-Fluorobiphenyl	15.52		19.47	0	79.7	70 - 130		
Surr: Trifluoromethyl benze				0	96.2	70 - 130		

Note: See Qualifiers Page for a list of qualifiers and their explanation.

Date: 28-Mar-18

Client: Columbia Technologies LLC

Project: Sac + Fox WorkOrder: HS18031138

QC BATCH REPORT

Batch ID: 126588	Inst	rument:	FID-10	Method: TX1005					
MSD Sample ID:	HS18031138-22MSD		Units: r	ng/Kg	Ana	alysis Date:	27-Mar-2018	11:53	П
Client ID: SB05-12	Run II	D: FID-1 0	_313250	SeqNo: 4	491758	PrepDate:	26-Mar-2018	DF: 1	
Analyte	Result	PQL	SPK Val	SPK Ref Value	%REC	Control Limit		RPD %RPD Limit	
nC6 to nC12	198.9	38	187.8	0	106	75 - 125	223.9	11.8 20)
>nC12 to nC28	195.8	38	187.8	0	104	75 - 125	209.2	6.6 20)
Surr: 2-Fluorobiphenyl	14.84	0	18.78	0	79.0	70 - 130	15.52	4.44 20)
Surr: Trifluoromethyl benzene	17.74	0	18.78	0	94.5	70 - 130	18.73	5.43 20)
The following samples were analyz	ed in this batch: HS180311 HS180311 HS180311 HS180311	138-16 138-20	HS18031138- HS18031138- HS18031138- HS18031138-	-17 -21	HS1803113 HS1803113 HS1803113 HS1803113	38-18 38-22	HS18031138- HS18031138- HS18031138-	19	1

Note: See Qualifiers Page for a list of qualifiers and their explanation.

Date: 28-Mar-18

Client: Columbia Technologies LLC

Serial Dilution

Sample Detection Limit

Texas Risk Reduction Program

Project: Sac + Fox

WorkOrder: HS18031138

SD

SDL

TRRP

QUALIFIERS, ACRONYMS, UNITS

-	
Qualifier	Description
*	Value exceeds Regulatory Limit
а	Not accredited
В	Analyte detected in the associated Method Blank above the Reporting Limit
E	Value above quantitation range
Н	Analyzed outside of Holding Time
J	Analyte detected below quantitation limit
M	Manually integrated, see raw data for justification
n	Not offered for accreditation
ND	Not Detected at the Reporting Limit
0	Sample amount is > 4 times amount spiked
Р	Dual Column results percent difference > 40%
R	RPD above laboratory control limit
S	Spike Recovery outside laboratory control limits
U	Analyzed but not detected above the MDL/SDL
Acronym	Description
DCS	Detectability Check Study
DUP	Method Duplicate
LCS	Laboratory Control Sample
LCSD	Laboratory Control Sample Duplicate
MBLK	Method Blank
MDL	Method Detection Limit
MQL	Method Quantitation Limit
MS	Matrix Spike
MSD	Matrix Spike Duplicate
PDS	Post Digestion Spike
PQL	Practical Quantitaion Limit

CERTIFICATIONS, ACCREDITATIONS & LICENSES

Date: 28-Mar-18

Agency	Number	Expire Date
California	2919 2016-2018	31-Jul-2018
Illinois	004112	09-May-2018
Kentucky	123043	30-Apr-2018
Louisiana	03087 2017-2017	30-Jun-2018
North Dakota	R193 2017-2017	30-Apr-2018
Oklahoma	2017-088	31-Aug-2018
Texas	T104704231-17-19	30-Apr-2018
North Carolina	624-2018	31-Dec-2018

Client: Columbia Technologies LLC

Project: Sac + Fox Work Order: HS18031138

SAMPLE TRACKING

Lab Samp ID	Client Sample ID	Action	Date	Person	New Location
HS18031138-01	SB01-11	Login	3/23/2018 8:16:19 PM	AV	LF021
HS18031138-02	SB01-12	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-03	SB01-13	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-04	SB01-14	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-05	SB01-15	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-06	SB02-12	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-07	SB02-14	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-08	SB02-16	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-09	SB03-20	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-10	SB03-21	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-11	SB03-22	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-12	SB04-11	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-13	SB04-12	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-14	SB04-13	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-15	SB04-14	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-16	SB04-15	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-17	SB04-16	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-18	SB05-8	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-19	SB05-9	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-20	SB05-10	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-21	SB05-11	Login	3/23/2018 8:25:53 PM	AV	LF021
HS18031138-22	SB05-12	Login	3/23/2018 8:25:54 PM	AV	LF021
HS18031138-23	SB05-13	Login	3/23/2018 8:25:54 PM	AV	LF021
HS18031138-24	SB05-14	Login	3/23/2018 8:25:54 PM	AV	LF021
HS18031138-25	SB05-15	Login	3/23/2018 8:25:54 PM	AV	LF021
HS18031138-26	SB05-16	Login	3/23/2018 8:25:54 PM	AV	LF021

Date: 28-Mar-18

Client Name: Vork Order:	ColumbiaTechnologies HS18031138			Fime Received: ved by:	Sample Rec 23-Mar-2018 PJM	ceipt Checklist 3 08:50
Checklist com	pleted by: Jared R. Makan eSignature	23-Mar-2018 Date	Reviewed by:	Corey Gra	ndits	26-Mar-2018 Date
Matrices:	Soil		Carrier name:	FedEx Price	ority Overnight	
Custody seals Custody seals Chain of custo Chain of custo Chain of custo Samples in pro Sample conta TX1005 solids Sufficient sam All samples re	dy signed when relinquished and receive dy agrees with sample labels? oper container/bottle?	Ye Y		No	Not Present Not Present Not Present	
	s)/Thermometer(s):	1.3	c/0.7c UC/C			IR25
Cooler(s)/Kit(s	s):	437	33			
	nple(s) sent to storage:	03/2	23/2018 20:40			
	rials have zero headspace? ceptable upon receipt? y:	Ye Ye	es 🗏	No No	N/A N/A	mitted
Login Notes:	Second vial for SB05-9' received cracke	ed, sample transferred	to new Tared vi	al, and flagged.		
Client Contact	ed: Da	ate Contacted:		Person Cor	ntacted:	
Contacted By:	Re	egarding:				
Comments:						
Corrective Act	ion:					



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COC ID: 180384

ALS Project Manager: ALS Work Order #: **Customer Information Project Information** Parameter/Method Request for Analysis Purchase Order Sac Project Name Pox TX1005_S_REV3 (5035/TPH TX1005 - *Freeze within 48 hours) Work Order В Project Number Company Name Bill To Company C Columbia Technologies LLC Columbia Technologies LLC Send Report To D Scott Pieper Invoice Attn Scott Pieper 1795 Cogswell Street 1795 Cogswell Street Е HS18031138 Address Address Suite 101 Suite 101 F Columbia Technologies LLC G City/State/Zip Rockledge,, NC 32955 City/State/Zip Rockledge, FL 32955 Sac + Fox Н Phone (410) 535-9911 Phone (410) 536-9911 Fax Fax SPieper@ColumbiaTechnologies.com SPieper@ColumbiaTechnologies.com e-Mail Address e-Mail Address No. Sample Description Date Time Matrix Pres. # Bottles Α В C D Ε G Н Hold SBOL -IL 1221 5:1 3/22 SB01-12 2 1226 5B01-13 2 2 Soi 1240 2 1306 1520 1326 2 ampler(s) Please Print & Sign Shipment Method Required Turnaround Time: (Check Box) Results Due Date: Other STD 10 Wk Days 5 Wk Days 2 Wk Davs 24 Hour Received by:
Received by (Laboratory): Date: 2 Notes: elinquished by: Time: Sutten ColumbiaTech - TPH 1600 Date: elinquished by:; Time: Cooler ID QC Package: (Check One Box Below) Fed Ex 1710 Mered Level II Std QC TRRP Checklist Checked by (Laboratory): Date: 43733 gged by (Laboratory): Time: PM 3-23-18 Level III Std QC/Raw Date TRRP Level IV 8 : 50 Level IV SW846/CLP eservative Key: 1-HCI 2-HNO₃ 3-H2SO4 4-NaOH 5-Na₂S₂O₃ 6-NaHSO 8-4°C 9-5035 7-Other

2: 1. Any changes must be made in writing once samples and COC Form have been submitted to ALS Environmental.

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	ALS Project Manager:								ALS	Work	Order	#:						
	Customer Information Project Inform				mati	on				Par	amete	r/Met	thod F	Reque	st for A	Analys	SIS	
Purchase Order		Project Nam	ne Sa	Sac + Fox				A	X1005	S_RE	EV3 (50)35/TI	PH TX	1005 -	*Free:	ze wit	hin 48 I	hours)
Work Order		Project Numb						В				7172 del 8/4					· · · · · · · · · · · · · · · · · · ·	A INTERNAL PROPERTY.
Company Name	Columbia Technologies LLC	Bill To Compar	עי Colu	mbia Te	echno	ologies LLC	2000	С		- Parkwall Illiands	2000							
Send Report To	Scott Pieper	Invoice At	tn Scot	t Pieper				D										
Address	1795 Cogswell Street Suite 101	Addres	1795 Cogswell Street					E	+ US18031138									
City/State/Zip	Rockledge,, NC 32955	City/State/Z	ip Rock	dedge, l	FL 3	2955		G		C	olum		i ecn Sac +		gies L	LC		
Phone	(410) 536-9911	Phor	ne (410) 536-99	911			Н										l
Fax		Fa	Fax				ı							A LEGISTRA				
e-Mail Address	SPieper@ColumbiaTechnologies.con	e-Mail Addres	e-Mail Address SPieper@ColumbiaTechnologies.com			J										i		
No.	Sample Description	Date	Time	Mati	rix	Pres.	# Bottles	Α	В	С	D	E	F	G	Н	ı	J	Hold
1 \$803-	22	3/22	1339	Soil			2											
2 SB04 -		_	1405	5011			2				78.					-		
3 5BOU -	12		1425	501			2											
4 SB04	- 13		1410	Sot			2				and the same							
5 SB04		3/22	1415	501			2											
6 5304 -	- 15		1420	501			2											
7 SB04 -		7/22	1430	50.			2								200			
8 5B05.		_	1510	501			2								VALUE			
9 SBO5-		3/22	1515	50,	· CORLOS		2											
10 SBO5-		3/22	1447	501	i		2				44						4	
Sampler(s) Please	Print & Sign ,	Shipment			Requ	uired Turnard	und Time: (Check	Box)	Oth	er			R	esults I	Due Da	te:	
Star 20		Fede			5	STD 10 Wk Day	s (Wk Da		2 \	k Days		24 F	lour		***********		No. of the last of
Relinquished by:	Aton 3/22	ime: 1600	Merce	lith	w)	latson	A CONTRACTOR OF THE PARTY OF TH	PERMINENTA	Col	THE PERSON NAMED IN	Tech - ler Temp.	examply terminated	Dooks	o. /Cha	k One B	ov Bolo		Andrewska on the William Court
Relinquished by:	Relinquished by: Agram Suffor Relinquished by: Date: Date: Time: Received by Received by Received by Time: Time: Received by Received by): P)M	AMOUNT AND	00	oler ID	1000	er lemp.		Level	III Std QC) D/Row Da		TRRE	P Checklist P Level IV
Preservative Key: 1-HCl 2-HNO ₃ 3-H ₂ SO ₄ 4-NaOH 5-Na ₂ S ₂ O ₃ 6-NaHSO ₄ 7-Other 8-4°C 9-5035																		

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		ALS Project Manager:															
Customer Information				ect Informat	tion		Parameter/Method Request for Analysis										
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Company Name	Columbia Technologies LLC	Bill To Compa	any Col	umbia Techi	nologies LL(>	С		~~~				w. water				A 15 A 15 A 16 A 16 A 16 A 16 A 16 A 16
Send Report To	Scott Pieper	Invoice A	ttn Sco	ott Pieper			D										
Address	1795 Cogswell Street Suite 101	Addre	226	1795 Cogswell Street Suite 101													
City/State/Zip	Rockledge,, NC 32955	City/State/2	Zip Ro	ckledge, FL	32955	444	G	Columbia Technologies LLC Sac + Fox									
Phone	(410) 536-9911	Pho	one (41	0) 536-9911			Н										
Fax		F	ax				1	-									
e-Mail Address	SPieper@ColumbiaTechnologies.com	e-Mail Addre	ess SPi	ieper@Colur	nbiaTechno	logies.com	J										
No.	Sample Description	Date	Time	Matrix	Pres.	# Bottles	Α	В	С	D	E	F	G	Н	I	J	Hold
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Preservative Key:	1-HCI 2-HNO ₃ 3-H ₂ SO ₄ 4-NaO	H 5-Na ₂ S ₂ O ₃	6-NaHS	3O ₄ 7-Othe	er 8-4°C	9-5035						-	Ē				

Any changes must be made in writing once samples and COC Form have been submitted to ALS Environmental.
 Unless otherwise agreed in a formal contract, services provided by ALS Environmental are expressly limited to the terms and conditions stated on the reverse.

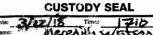
3. The Chain of Custody is a legal document. All information must be completed accurately.

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teranext



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CUSTODY SEAL

Date: 3/72//8 Time: 17-10
Name: Mered H Wetson
Company: Terrenext

3-23-16

TRK# 7376 9753 4247

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List of Symbols, Abbreviations, and Acronyms

CSM Conceptual Site Model. A CSM is a method to describe what is

known or can be inferred about a site for the purpose of making a decision. A CSM generally will address physical, chemical and biological systems; contaminant release and transport; societal

issues; policy, land use, and exposures.

CVOC Chlorinated Volatile Organic Contaminant. A VOC containing

chlorine atoms; typically, a cleaning solvent.

DPT Direct-Push Technology (DPT) refers to a group of techniques

used for subsurface investigation by driving, pushing and/or

vibrating small-diameter rods into the ground.

ECD Electron Capture Detector. An ECD is a device for detecting

electron-absorbing components (high electronegativity) such as halogenated compounds in a gas through the attachment of

electrons via electron capture ionization.

DNAPL Dense Non-Aqueous Phase Liquid. A DNAPL is a denser-than-

water NAPL, i.e. a liquid that is both denser than water and is

immiscible in or does not dissolve in water.

HPT Hydraulic Profiling Tool. The HPT is a logging tool that

measures the pressure required to inject a flow of water into the soil as the probe is advanced into the subsurface. In addition to measurement of injection pressure, the HPT can also be used to measure hydrostatic pressure under the zero flow condition.

LCSM LNAPL Conceptual Site Model. A LCSM is a conceptual site

model focused on the release and transport of LNAPL

contaminants.

LIF Laser-induced fluorescence is a spectroscopic method in which

an atom or molecule is excited to a higher energy level by the absorption of laser light followed by spontaneous emission of light.

LNAPL Light Non-Aqueous Phase Liquids are groundwater

contaminants that are not soluble in water and have lower density than water, in contrast to a **DNAPL** which has higher density than

water.

Symbol or Abbreviation	Definition
PHC	Petroleum Hydrocarbons. The presence of petroleum hydrocarbon fuels in any phase. (PHC).
PID	Photo Ionization Detector. In a PID high-energy photons to break molecules into positively charged ions. The PID will only respond to components that have ionization energies at or below the energy of the photons produced by the PID lamp.
SPOC	Shock Protected Optical Cavity . The SPOC is the component of the LIF system that contains the mirror and sapphire window for proper alignment of the laser beam.
TCE	Trichloroethylene. The chemical compound TCE is a halocarbon commonly used as an industrial solvent. It is a clear non-flammable liquid with a sweet smell.
UST	Underground Storage Tank. Under Federal law UST means any one or combination of tanks including connected underground pipes that is used to contain regulated substances, and the volume of which including the volume of underground pipes is 10 percent or more beneath the surface of the ground. This does not include, among other things, any farm or residential tank of 1,100 gallons or less capacity used for storing motor fuel for noncommercial purposes, tanks for storing heating oil for consumption on the premises, or septic tanks.
UVOST®	Ultraviolet Optical Scanning Tool ®. A LIF is a tool that uses laser light in the ultraviolet spectrum to excite fluorescent molecules that exist in the vast majority of hazardous non-aqueous phase liquids (NAPLs) such as petroleum fuels/oils, coal tars, and creosotes.
VOC	Volatile organic compounds (VOCs) are organic chemicals that have a high vapor pressure at ordinary room temperature. Their high vapor pressure results from a low boiling point, which causes large numbers of molecules to evaporate or sublimate from the liquid or solid form of the compound and enter the surrounding air, a trait known as volatility.
XSD	Halogen Specific Detector. The XSD was developed for the selective detection of halogen-containing compounds.

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